

(19)



Europäisches Patentamt
European Patent Office
Office européen des brevets

(11) Publication number:

**0 325 262
A2**

(12)

EUROPEAN PATENT APPLICATION

(21) Application number: 89100913.6

(51) Int. Cl.4: **C12N 15/00 , C12P 21/02 ,
A61K 39/395 , G01N 33/569 ,
//A61K39/21**

(22) Date of filing: 20.01.89

The applicant has filed a statement in accordance with Rule 28 (4) EPC (issue of a sample only to an expert). Accession number(s) of the deposit(s): ATCC 67608 - 67609 - 67610 - 67611.

Claims for the following Contracting States: ES
+ GR.

(30) Priority: 22.01.88 US 147351

(43) Date of publication of application:
26.07.89 Bulletin 89/30

(84) Designated Contracting States:
AT BE CH DE ES FR GB GR IT LI LU NL SE

(71) Applicant: **THE GENERAL HOSPITAL
CORPORATION**
55 Fruit Street
Boston MA 02114(US)

(72) Inventor: **Brian, Seed, Dr.**
47A Joy Street
Boston MA 02114(US)

(74) Representative: **Klein, Otto, Dr. et al**
Hoechst AG Zentrale Patentabteilung
Postfach 80 03 20
D-6230 Frankfurt am Main 80(DE)

(54) **Cloned genes encoding IG-CD4 fusion proteins and the use thereof.**

(57) Fusion proteins of immunoglobulins of the IgM, IgG1 or IgG3 class, wherein the variable region of the light or heavy chain has been replaced with CD4 or fragments thereof capable of binding to gp120 or immunoglobulin-like molecules comprising such fusion proteins together with an immunoglobulin light or heavy chain can be administered to an animal suffering from HIV or SIV infection. They also are useful in assays for HIV or SIV comprising contacting a sample suspected of containing HIV or SIV gp120 with the immunoglobulin-like molecule or fusion protein, and detecting whether a complex is formed.

EP 0 325 262 A2

CLONED GENES ENCODING IG-CD4 FUSION PROTEINS AND THE USE THEREOF

CROSS-REFERENCE TO RELATED APPLICATION

This application is a continuation-in-part of U.S. Application Serial No. 07/147,351 filed January 22, 1988.

FIELD OF THE INVENTION

The invention is in the field of recombinant genetics.

BACKGROUND OF THE INVENTION

The human and simian immunodeficiency viruses HIV and SIV are the causative agents of Acquired Immune Deficiency Syndrome (AIDS) and Simian Immunodeficiency Syndrome (SIDS), respectively. See Curren, J. et al., *Science* 329:1359-1357 (1985); Weiss, R. et al., *Nature* 324:572-575 (1986). The HIV virus contains an envelope glycoprotein, gp120 which binds to the CD4 protein present on the surface of helper T lymphocytes, macrophages and other cells. Dalglish et al., *Nature*, 312:763 (1984). After the gp120 binds to CD4, virus entry is facilitated by an envelope-mediated fusion of the viral target cell membranes.

During the course of infection, the host organism develops antibodies against viral proteins, including the major envelope glycoproteins gp120 and gp41. Despite this humoral immunity, the disease progresses, resulting in a lethal immunosuppression characterized by multiple opportunistic infections, parasitemia, dementia and death. The failure of host anti-viral antibodies to arrest the progression of the disease represents one of the most vexing and alarming aspects of the infection, and augurs poorly for vaccination efforts based upon conventional approaches.

Two factors may play a role in the inefficacy of the humoral response to immunodeficiency viruses. First, like other RNA viruses (and like retroviruses in particular), the immunodeficiency viruses show a high mutation rate which allows antigenic variation to progress at a high rate in response to host immune surveillance. Second, the envelope glycoproteins themselves are heavily glycosylated molecules presenting few epitopes suitable for high affinity antibody binding. The poorly antigenic, "moving" target which the viral envelope presents, allows the host little opportunity for restricting viral infection by specific antibody production.

Cells infected by the HIV virus express the gp120 glycoprotein on their surface. Gp120 mediates fusion events among CD4⁺ cells via a reaction similar to that by which the virus enters the uninfected cell, leading to the formation of short-lived multinucleated giant cells. Syncytium formation is dependent on a direct interaction of the gp120 envelope glycoprotein with the CD4 protein. Dalglish et al., *supra*, Klatzmann, D. et al., *Nature* 312:763 (1984); McDougal, J.S. et al., *Science*, 231:382 (1986); Sodroski, J. et al., *Nature*, 322:470 (1986); Lifson, J.D. et al., *Nature*, 323:725 (1986); Sodroski, J. et al., *Nature*, 321:412 (1986).

The CD4 protein consists of a 370 amino acid extracellular region containing four immunoglobulin-like domains, a membrane spanning domain, and a charged intracellular region of 40 amino acid residues. Maddon, P. et al., *Cell* 42:93 (1985); Clark, S. et al., *Proc. Natl. Acad. Sci. (USA)* 84:1649 (1987).

Evidence that CD4-gp120 binding is responsible for viral infection of cells bearing the CD4 antigen includes the finding that a specific complex is formed between gp120 and CD4. McDougal et al., *supra*. Other workers have shown that cell lines, which were non-infective for HIV, were converted to infectable cell lines following transfection and expression of the human CD4 cDNA gene. Maddon et al., *Cell* 47:333-348 (1986).

In contrast to the majority of antibody-envelope interactions, the receptor-envelope interaction is characterized by a high affinity ($K_a \approx 10^8$ /mole) immutable association. Moreover, the affinity of the virus for CD4 is at least 3 orders of magnitude higher than the affinity of CD4 for its putative endogenous ligand, the MHC class II antigens. Indeed, to date, a specific physical association between monomeric CD4 and class II antigens has not been demonstrated.

In response to bacterial or other particle infection, the host organism usually produces serum antibodies that bind to specific proteins or carbohydrates on the bacterial or particle surface, coating the bacteria. This antibody coat on the bacterium or other particle stimulates cytotoxicity by Fc-receptor-bearing lymphoid cells by antibody-dependent cellular toxicity (ADCC). Other serum proteins, collectively called complement (C),

bind to antibody-coated targets, and also can coat foreign particles nonspecifically. They cause cell death by lysis, or stimulate ingestion by binding to specific receptors on the macrophage called complement receptors. See Darnell J. et al., in Molecular Cell Biology, Scientific American Books, pp. 641 and 1087 (1986).

5 The most effective complement activating classes of human Ig are IgM and IgG1. The complement system consists of 14 proteins that, acting in order, cause lysis of cells. Nearly all of the C proteins exist in normal serum as inactive precursors. When activated, some become highly specific proteolytic enzymes whose substrate is the next protein in a sequential chain reaction.

The entire C sequence can be triggered by either of two initiation pathways. In one (the classic 10 pathway), Ab-Ag complexes bind and activate C1, C4 and C2 to form a C3-splitting enzyme. In the second pathway, polysaccharides commonly on the surface of many bacteria and fungi bind with trace amounts of a C3 fragment and then with two other proteins (factor B and properdin) to form another C3-splitting enzyme. Once C3 is split by either pathway, the way is open for the remaining sequence of steps which lead to cell lysis. See Davis, B.D., et al., in Microbiology, 3rd ed., Harper and Row, Philadelphia, PA, pp. 15 452-466 (1980).

A number of workers have disclosed methods for preparing hybrid proteins. For example, Murphy, United States Patent 4,675,382 (1987), discloses the use of recombinant DNA techniques to make hybrid protein molecules by forming the desired fused gene coding for a hybrid protein of diphtheria toxin and a polypeptide ligand such as a hormone, followed by expression of the fused gene.

20 Many workers have prepared monoclonal antibodies (Mabs) by recombinant DNA techniques. Monoclonal antibodies are highly specific well-characterized molecules in both primary and tertiary structure. They have been widely used for in vitro immunochemical characterization and quantitation of antigens. Genes for heavy and light chains have been introduced into appropriate hosts and expressed, followed by reaggregation of the individual chains into functional antibody molecules (see, for example, Munro, Nature 25 312:597 (1984); Morrison, S.L., Science 229:1202 (1985); Oi et al., Biotechniques 4:214 (1986); Wood et al., Nature 314:446-449 (1985)). Light- and heavy-chain variable regions have been cloned and expressed in foreign hosts wherein they maintained their binding ability (Moore et al., European Patent Application 0088994 (published September 21, 1983)).

Chimeric or hybrid antibodies have also been prepared by recombinant DNA techniques. Oi and 30 Morrison, Biotechniques 4:214 (1986) describe a strategy for producing such chimeric antibodies which include a chimeric human IgG anti-leu3 antibody.

Gascoigne, N.R.J., et al., Proc. Natl. Acad. Sci. (USA) 84:2936-2940 (1987) disclose the preparation of a chimeric gene construct containing a T-cell receptor α -chain variable (V) domain and the constant (C) region coding sequence of an immunoglobulin γ 2a molecule. Cells transfected with the chimeric gene synthesize a 35 protein product that expresses immunoglobulin and T-cell receptor antigenic determinants as well as protein A binding sites. This protein associates with a normal λ chain to form an apparently normal tetrameric (H_2L_2 , where H = heavy and L = light) immunoglobulin molecule that is secreted.

Sharon, J., et al., Nature 309:54 (1984), disclose construction of a chimeric gene encoding the variable (V) region of a mouse heavy chain specific for the hapten azophenylarsonate and the constant (C) region of 40 a mouse kappa light chain ($V_H C_K$). This gene was introduced into a mouse myeloma cell line. The chimeric gene was expressed to give a protein which associated with light chains secreted from the myeloma cell line to give an antibody molecule specific for azophenylarsonate.

Morrison, Science 229:1202 (1985), discloses that variable light-or variable heavy-chain regions can be attached to a non-Ig sequence to create fusion proteins. This article states that the potential uses for the 45 fusion proteins are three: (1) to attach antibody specifically to enzymes for use in assays; (2) to isolate non-Ig proteins by antigen columns; and (3) to specifically deliver toxic agents.

Recent techniques for the stable introduction of immunoglobulin genes into myeloma cells (Banerji, J., et al., Cell 33:729-740 (1983); Potter, H., et al., Proc. Natl. Acad. Sci. (USA) 81:7161-7165 (1984)), coupled 50 with detailed structural information, have permitted the use of in vitro DNA methods such as mutagenesis, to generate recombinant antibodies possessing novel properties.

PCT Application W087/02671 discloses methods for producing genetically engineered antibodies of desired variable region specificity and constant region properties through gene cloning and expression of light and heavy chains. The mRNA from cloned hybridoma B cell lines which produce monoclonal antibodies of desired specificity is isolated for cDNA cloning. The generation of light and heavy chain 55 coding sequences is accomplished by excising the cloned variable regions and ligating them to light or heavy chain module vectors. This gives cDNA sequences which code for immunoglobulin chains. The lack of introns allows these cDNA sequences to be expressed in prokaryotic hosts, such as bacteria, or in lower eukaryotic hosts, such as yeast.

The generation of chimeric antibodies in which the antigen-binding portion of the immunoglobulin is fused to other moieties has been demonstrated. Examples of non-immunoglobulin genes fused to antibodies include *Staphylococcus aureus* nuclease, the mouse oncogene c-myc, and the Klenow fragment of *E. coli* DNA polymerase I (Neuberger, M.S., et al., *Nature* 312:604-612 (1984); Neuberger, M.S., *Trends in Biochemical Science*, 347-349 (1985)). European Patent Application 120,694 discloses the genetic engineering of the variable and constant regions of an immunoglobulin molecule that is expressed in *E. coli* host cells. It is further disclosed that the immunoglobulin molecule may be synthesized by a host cell with another peptide moiety attached to one of the constant domains. Such peptide moieties are described as either cytotoxic or enzymatic. The application and the examples describe the use of a lambda-like chain derived from a monoclonal antibody which binds to 4-hydroxy-3-nitrophenyl (NP) haptens.

European Patent Application 125,023 relates to the use of recombinant DNA techniques to produce immunoglobulin molecules that are chimeric or otherwise modified. One of the uses described for these immunoglobulin molecules is for whole-body diagnosis and treatment by injection of the antibodies directed to specific target tissues. The presence of the disease can be determined by attaching a suitable label to the antibodies, or the diseased tissue can be attacked by carrying a suitable drug with the antibodies. The application describes antibodies engineered to aid the specific delivery of an agent as "altered antibodies."

PCT Application W083/101533 describes chimeric antibodies wherein the variable region of an immunoglobulin molecule is linked to a portion of a second protein which may comprise the active portion of an enzyme.

Boulianne et al., *Nature* 312:643 (1984) constructed an immunoglobulin gene in which the DNA segments that encode mouse variable regions specific for the hapten trinitrophenol (TNP) are joined to segments that encode human mu and kappa regions. These chimeric genes were expressed to give functional TNP-binding chimeric IgM.

Morrison et al., *P.N.A.S. (USA)* 81:6851 (1984), disclose a chimeric molecule utilizing the heavy-chain variable region exons of an anti-phosphoryl choline myeloma protein G, which were joined to the exons of either human kappa light-chain gene. The genes were transfected into mouse myeloma cell lines, generating transformed cells that produced chimeric mouse-human IgG with antigen-binding function.

Despite the progress that has been achieved on determining the mechanism of HIV infection, a need continues to exist for methods of treating HIV viral infections.

SUMMARY OF THE INVENTION

The invention relates to a gene comprising a DNA sequence which encodes a fusion protein comprising 1) CD4, or a fragment thereof which binds to HIV gp120, and 2) an immunoglobulin light or heavy chain; wherein said CD4 or HIV gp120-binding fragment thereof replaces the variable region of the light or heavy immunoglobulin chain.

The invention also relates to vectors containing the gene of the invention and hosts transformed with the vectors.

The invention also relates to a method of producing a fusion protein comprising CD4, or fragment thereof which binds to HIV gp120, and an immunoglobulin light or heavy chain, wherein the variable region of the immunoglobulin light or heavy chain has been substituted with CD4, or HIV gp120-binding fragment thereof, which comprises:

cultivating in a nutrient medium under protein producing conditions, a host strain transformed with the vector containing the gene of the invention, said vector further comprising expression signals which are recognized by said host strain and direct expression of said fusion protein, and recovering the fusion protein so produced.

The invention also relates to a fusion protein comprising CD4, or fragment thereof which is capable of binding to HIV gp120, fused at the C-terminus to a second protein which comprises an immunoglobulin light or heavy chain, wherein the variable region of said light or heavy chain is substituted with CD4 or a HIV gp120 binding fragment thereof.

The invention also relates to an immunoglobulin-like molecule comprising the fusion protein of the invention together with an immunoglobulin light or heavy-chain, wherein said immunoglobulin like molecule binds HIV gp120.

The IgG1 fusion proteins and immunoglobulin-like molecules may be useful for both complement-mediated and cell-mediated (ADCC) immunity, while the IgM fusion proteins are useful principally through complement-mediated immunity.

The invention also relates to a complex between the fusion proteins and immunoglobulin-like molecule of the invention and HIV gp120.

The invention also relates to a method for treating HIV or SIV infections comprising administering the fusion protein or immunoglobulin-like molecule of the invention to an animal.

5 The invention further relates to a method for detecting HIV gp120 in a sample comprising contacting a sample suspected of containing HIV or gp120 with the fusion protein or immunoglobulin-like molecule of the invention, and detecting whether a complex has formed.

10 DESCRIPTION OF THE PREFERRED EMBODIMENTS

The invention is directed to a protein gene which comprises

- 1) a DNA sequence which codes for CD4, or fragment thereof which binds to HIV gp120, fused to
- 15 2) a DNA sequence which encodes an immunoglobulin heavy chain.

Preferably, the antibody has effector function.

The invention is also directed to a protein gene which comprises

- 1) a DNA sequence which codes for CD4, or fragment thereof which binds to HIV gp120, fused to
- 20 2) a DNA sequence which encodes an immunoglobulin light chain; wherein said sequence which codes for CD4, or HIV gp120-binding fragment thereof, replaces the variable region of the light immunoglobulin chain.

The invention is also directed to the expression of these novel fusion proteins in transformed hosts and the use thereof to treat and diagnose HIV infections. In particular, the invention relates to expressing said genes in mammalian hosts which express complementary light or heavy chain immunoglobulins to give immunoglobulin-like molecules which have antibody effector function and also bind to HIV or SIV gp120.

The term "antibody effector function" as used herein denotes the ability to fix complement or to activate ADCC.

30 The fusion proteins and immunoglobulin-like molecules may be administered to an animal for the purpose of treating HIV or SIV infections. By the terms "HIV infections" is intended the condition of having AIDS, AIDS related complex (ARC) or where an animal harbors the AIDS virus, but does not exhibit the clinical symptoms of AIDS or ARC. By the terms "SIV infections" is intended the condition of being infected with simian immunodeficiency virus.

35 By the term "animal" is intended all animals which may derive benefit from the administration of the fusion proteins and immunoglobulin-like molecules of the invention. Foremost among such animals are humans, however, the invention is not intended to be so limited.

By the term "fusion protein" is intended a fused protein comprising CD4, or fragment thereof which is capable of binding to gp120, linked at its C-terminus to an immunoglobulin chain wherein a portion of the N-terminus of the immunoglobulin is replaced with CD4. In general, that portion of immunoglobulin which is 40 deleted is the variable region. The fusion proteins of the invention may also comprise immunoglobulins where more than just the variable region has been deleted and replaced with CD4 or HIV gp120 binding fragment thereof. For example, the V_H and CH1 regions of an immunoglobulin chain may be deleted. Preferably, any amount of the N-terminus of the immunoglobulin heavy chain can be deleted as long as the remaining fragment has antibody effector function. The minimum sequence required for binding complement encompasses domains CH2 and CH3. Joining of Fc portions by the hinge region is advantageous for increasing the efficiency of complement binding.

The CD4 portion of the fusion protein may comprise the complete CD4 sequence, the 370 amino acid extracellular region and the membrane spanning domain, or the extracellular region. The fusion protein may 50 comprise fragments of the extracellular region obtained by cutting the DNA sequence which encodes CD4 at the BspM1 site at position 514 or the PvuII site at position 629 (see Table 1) to give nucleotide sequences which encode CD4 fragments which retain binding to gp120. In general, any fragment of CD4 may be used as long as it retains binding to gp120.

Where the fusion protein comprises an immunoglobulin light chain, it is necessary that no more of the 55 Ig chain be deleted than is necessary to form a stable complex with a heavy chain Ig. In particular, the cysteine residues necessary for disulfide bond formation must be preserved on both the heavy and light chain moieties.

When expressed in a host, e.g., a mammalian cell, the fusion protein may associate with other light or

heavy Ig chains secreted by the cell to give a functioning immunoglobulin-like molecule which is capable of binding to gp120. The gp120 may be in solution, expressed on the surface of infected cells, or may be present on the surface of the HIV virus itself. Alternatively, the fusion protein may be expressed in a mammalian cell which does not secrete other light or heavy Ig chains. When expressed under these conditions, the fusion protein may form a homodimer.

Genomic or cDNA sequences may be used in the practice of the invention. Genomic sequences are expressed efficiently in myeloma cells, since they contain native promoter structures.

The constant regions of the antibody cloned and used in the chimeric immunoglobulin-like molecule may be derived from any mammalian source. The constant regions may be complement binding or ADCC active. However, preliminary work (see Examples) indicates that the fusion proteins of the invention may mediate HIV or SIV infected cell death by an ADCC or complement-independent mechanism. The constant regions may be derived from any appropriate isotype, including IgG1, IgG3, or IgM.

The joining of various DNA fragments, is performed in accordance with conventional techniques, employing blunt-ended or staggered-ended termini for ligation, restriction enzyme digestion to provide appropriate termini, filling in of cohesive ends as appropriate, alkali and phosphatase treatment to avoid undesirable joining, and ligation with appropriate ligases. The genetic construct may optionally encode a leader sequence to allow efficient expression of the fusion protein. For example, the leader sequence utilized by Maddon et al., *Cell* 42:93-104 (1985) for the expression of CD4 may be used.

For cDNA, the cDNA may be cloned and the resulting clone screened, for example, by use of a complementary probe or by assay for expressed CD4 using an antibody as disclosed by Dalglish et al., *Nature* 312:763-766 (1984); Klatzmann et al., *Immunol. Today* 7:291-297 (1986); McDougal et al., *J. Immunol.* 135:3151-3162 (1985); and McDougal, J. et al., *J. Immunol.* 137:2937-2944 (1986).

To express the fusion hybrid protein, transcriptional and translational signals recognized by an appropriate host element are necessary. Eukaryotic hosts which may be used include mammalian cells capable of culture in vitro, particularly leukocytes, more particularly myeloma cells or other transformed or oncogenic lymphocytes, e.g., EBV-transformed cells. Alternatively, non-mammalian cells may be employed, such as bacteria, fungi, e.g., yeast, filamentous fungi, or the like.

Preferred hosts for fusion protein production are mammalian cells, grown in vitro in tissue culture or in vivo in animals. Mammalian cells provide post translational modification to immunoglobulin protein molecules which provide for correct folding and glycosylation of appropriate sites. Mammalian cells which may be useful as hosts include cells of fibroblast origins such as VERO or CHO-K1 or cells of lymphoid origin, such as the hybridoma SP2/0-AG14 or the myeloma P3x63Sgh, and their derivatives. For the purpose of preparing an immunoglobulin-like molecule, a plasmid containing a gene which encodes a heavy chain immunoglobulin, wherein the variable region has been replaced with CD4 or fragment thereof which binds to gp120, may be introduced, for example, into J558L myeloma cells, a mouse plasmacytoma expressing the lambda-1 light chain but which does not express a heavy chain (see Oi et al., *P.N.A.S. (USA)* 80:825-829 (1983)). Other preferred hosts include COS cells, BHK cells and hepatoma cells.

The constructs may be joined together to form a single DNA segment or may be maintained as separate segments, by themselves or in conjunction with vectors.

Where the fusion protein is not glycosylated, any host may be used to express the protein which is compatible with replicon and control sequences in the expression plasmid. In general, vectors containing replicon and control sequences are derived from species compatible with a host cell are used in connection with the host. The vector ordinarily carries a replicon site, as well as specific genes which are capable of providing phenotypic selection in transformed cells. The expression of the fusion protein can also be placed under control with other regulatory sequences which may be homologous to the organism in its untransformed state. For example, lactose-dependent *E. coli* chromosomal DNA comprises a lactose or lac operon which mediates lactose utilization by elaborating the enzyme beta-galactosidase. The lac control elements may be obtained from bacterial phage lambda plac5, which is infective for *E. coli*. The lac promoter-operator system can be induced by IPTG.

Other promoters/operator systems or portions thereof can be employed as well. For example, colicin E1, galactose, alkaline phosphatase, tryptophan, xylose, tax, and the like can be used.

For mammalian hosts, several possible vector systems are available for expression. One class of vectors utilize DNA elements which are derived from animal viruses such as bovine papilloma virus, polyoma virus, adenovirus, vaccinia virus, baculovirus, retroviruses RSV, MMTV or MOMLV), or SV40 virus. Cells which have stably integrated the DNA into their chromosomes may be selected by introducing one or more markers which allow selection of transfected host cells. The marker may provide for prototrophy to an auxotrophic host, biocide resistance, e.g., antibiotics, or heavy metals such as copper or the like. The selectable marker gene can be either directly linked to the DNA sequences to be expressed, or introduced

into the same cell by cotransformation. Additional elements may also be needed for optimal synthesis of mRNA. These elements may include splice signals, as well as transcriptional promoters, enhancers, and termination signals. The cDNA expression vectors incorporating such elements includes those described by Okayama, H., Mol. Cel. Biol., 3:280 (1983) and others.

5 Once the vector or DNA sequence containing the constructs has been prepared for expression, the DNA constructs may be introduced to an appropriate host. Various techniques may be employed, such as protoplast fusion, calcium phosphate precipitation, electroporation or other conventional techniques. After the fusion, the cells are grown in media and screened for the appropriate activity. Expression of the gene(s) results in production of the fusion protein. This expressed fusion protein may then be subject to further
10 assembly to form the immunoglobulin-like molecule.

The host cells for immunoglobulin production may be immortalized cells, primarily myeloma or lymphoma cells. These cells may be grown in appropriate nutrient medium in culture flasks or injected into a synergistic host, e.g., mouse or a rat, or immunodeficient host or host site, e.g., nude mouse or hamster pouch. In particular, the cells may be introduced into the abdominal cavity of an animal to allow production
15 of ascites fluid which contains the immunoglobulin-like molecule. Alternatively, the cells may be injected subcutaneously and the chimeric antibody is harvested from the blood of the host. The cells may be used in the same manner as hybridoma cells. See Diamond et al., N. Eng. J. Med. 304:1344 (1981), and Kennatt, McKearn and Bechtol (Eds.), Monoclonal Antibodies: Hybridomas: -- A New Dimension in Biologic Analysis, Plenum, 1980.

20 The fusion proteins and immunoglobulin-like molecules of the invention may be isolated and purified in accordance with conventional conditions, such as extraction, precipitation, chromatography, affinity chromatography, electrophoresis or the like. For example, the IgG1 fusion proteins may be purified by passing a solution through a column which contains immobilized protein A or protein G which selectively binds the Fc portion of the fusion protein. See, for example, Reis, K.J., et al., J. Immunol. 132:3098-3102
25 (1984); PCT Application, Publication No. W087/00329. The chimeric antibody may be eluted by treatment with a chaotropic salt or by elution with aqueous acetic acid (1 M).

Alternatively the fusion proteins may be purified on anti-CD4 antibody columns, or on anti-immunoglobulin antibody columns.

In one embodiment of the invention, cDNA sequences which encode CD4, or a fragment thereof which
30 binds gp120, may be ligated into an expression plasmid which codes for an antibody wherein the variable region of the gene has been deleted. Methods for the preparation of genes which encode the heavy or light chain constant regions of immunoglobulins are taught, for example, by Robinson, R. et al., PCT Application, Publication No. W087-02671.

Preferred immunoglobulin-like molecules which contain CD4, or fragments thereof, contain the constant
35 region of an IgM, IgG1 or IgG3 antibody which binds complement at the Fc region.

The fusion protein and immunoglobulin-like molecules of the invention may be used for the treatment of HIV viral infections. The fusion protein complexes to gp120 which is expressed on infected cells. Although the inventor is not bound by a particular theory, it appears that the Fc portion of the hybrid fusion protein may bind with complement, which mediates destruction of the cell. In this manner, infected cells are
40 destroyed so that additional viral particle production is stopped.

For the purpose of treating HIV infections, the fusion protein or immunoglobulin-like molecule of the invention may additionally contain a radiolabel or therapeutic agent which enhances destruction of the HIV particle or HIV-infected cell.

Examples of radioisotopes which can be bound to the fusion protein or immunoglobulin-like molecule of
45 the invention for use in HIV-therapy are ^{125}I , ^{131}I , ^{90}Y , ^{67}Cu , ^{217}Bi , ^{211}At , ^{212}Pb , ^{47}Sc , and ^{109}Pd . Optionally, a label such as boron can be used which emits α and β particles upon bombardment with neutron radiation.

For in vivo diagnosis radionucleotides may be bound to the fusion protein or immunoglobulin-like molecule of the invention either directly or by using an intermediary functional group. An intermediary group which is often used to bind radioisotopes, which exist as metallic cations, to antibodies is
50 diethylenetriaminepentaacetic acid (DTPA). Typical examples of metallic cations which are bound in this manner are $^{99\text{m}}\text{Tc}$, ^{123}I , ^{111}In , ^{131}I , ^{97}Ru , ^{67}Cu , ^{67}Ga , and ^{68}Ga .

Moreover, the fusion protein and immunoglobulin-like molecule of the invention may be tagged with an NMR imaging agent which include paramagnetic atoms. The use of an NMR imaging agent allows the in vivo diagnosis of the presence of and the extent of HIV infection within a patient using NMR techniques.
55 Elements which are particularly useful in this manner are ^{157}Gd , ^{55}Mn , ^{162}Dy , ^{52}Cr , and ^{56}Fe .

Therapeutic agents may include, for example, bacterial toxins such as diphtheria toxin, or ricin. Methods for producing fusion proteins comprising fragment A of diphtheria toxin are taught in U.S. Patent 4,675,382 (1987). Diphtheria toxin contains two polypeptide chains. The B chain binds the toxin to a receptor on a cell

surface. The A chain actually enters the cytoplasm and inhibits protein synthesis by inactivating elongation factor 2, the factor that translocates ribosomes along mRNA concomitant with hydrolysis of ETP. See Darnell, J., et al., in Molecular Cell Biology, Scientific American Books, Inc., page 662 (1986). Alternatively, a fusion protein comprising ricin, a toxic lectin, may be prepared.

5 Introduction of the chimeric molecules by gene therapy may also be contemplated, for example, using retroviruses or other means to introduce the genetic material encoding the fusion proteins into suitable target tissues. In this embodiment, the target tissues having the cloned genes of the invention may then produce the fusion protein in vivo.

10 The dose ranges for the administration of the fusion protein or immunoglobulin-like molecule of the invention are those which are large enough to produce the desired effect whereby the symptoms of HIV or SIV infection are ameliorated. The dosage should not be so large as to cause adverse side effects, such as unwanted cross-reactions, anaphylactic reactions, and the like. Generally, the dosage will vary with the age, condition, sex and extent of disease in the patient, counterindications, if any, immune tolerance and other such variables, to be adjusted by the individual physician. Dosage can vary from .01 mg/kg to 50 mg/kg, 15 preferably 0.1 mg/kg to 1.0 mg/kg, of the immunoglobulin-like molecule in one or more administrations daily, for one or several days. The immunoglobulin-like molecule can be administered parenterally by injection or by gradual perfusion over time. They can be administered intravenously, intraperitoneally, intramuscularly, or subcutaneously.

20 Preparations for parenteral administration include sterile or aqueous or non-aqueous solutions, suspensions, and emulsions. Examples of non-aqueous solvents are propylene glycol, polyethylene glycol, vegetable oils such as olive oil, and injectable organic esters such as ethyl oleate. Aqueous carriers include water, alcoholic/aqueous solutions, emulsions or suspensions, including saline and buffered media. Parenteral vehicles include sodium chloride solution, Ringer's dextrose, dextrose and sodium chloride, lactated Ringer's, or fixed oils. Intravenous vehicles include fluid and nutrient replenishers, electrolyte replenishers, 25 such as those based on Ringer's dextrose, and the like. Preservatives and other additives may also be present, such as, for example, antimicrobials, antioxidants, chelating agents, inert gases and the like. See, generally, Remington's Pharmaceutical Science, 16th Ed., Mack Eds., 1980.

30 The invention also relates to a method for preparing a medicament or pharmaceutical composition comprising the components of the invention, the medicament being used for therapy of HIV or SIV infection in animals.

The detection and quantitation of antigenic substances and biological samples frequently utilized immunoassay techniques. These techniques are based upon the formation of the complex between the antigenic substance, e.g., gp120, being assayed and an antibody or antibodies in which one or the other member of the complex may be detectably labeled. In the present invention, the immunoglobulin-like 35 molecule or fusion protein may be labeled with any conventional label.

Thus, the hybrid fusion protein or immunoglobulin-like molecule of the invention can also be used in assay for HIV or SIV viral infection in a biological sample by contacting a sample, derived from an animal suspected of having an HIV or SIV infection, with the fusion protein or immunoglobulin-like molecule of the invention, and detecting whether a complex with gp120, either alone or on the surface of an HIV-infected 40 cell, has formed.

For example, a biological sample may be treated with nitrocellulose, or other solid support which is capable of immobilizing cells, cell particles or soluble protein. The support may then be washed with suitable buffers followed by treatment with the fusion protein which may be detectably labeled. The solid phase support may then be washed with the buffer a second time to remove unbound fusion protein and 45 the label on the fusion protein detected.

In carrying out the assay of the present invention on a sample containing gp120, the process comprises:

- a) contacting a sample suspected containing gp120 with a solid support to effect immobilization of gp120, or cell which expresses gp120 on its surface;
- 50 b) contacting said solid support with the detectably labeled immunoglobulin-like molecule or fusion protein of the invention;
- c) incubating said detectably labeled immunoglobulin-like molecule with said support for a sufficient amount of time to allow the immunoglobulin-like molecule or fusion protein to bind to the immobilized gp120 or cell which expresses gp120 on its surface;
- 55 d) separating the solid phase support from the incubation mixture obtained in step c); and
- e) detecting the bound immunoglobulin-like molecule or fusion protein and thereby detecting and quantifying gp120.

Alternatively, labeled immunoglobulin-like molecule (or fusion protein) -gp120 complex in a sample may be separated from a reaction mixture by contacting the complex with an immobilized antibody or protein which is specific for an immunoglobulin or, e.g., protein A, protein G, anti-IgM or anti-IgG antibodies. Such anti-immunoglobulin antibodies may be monoclonal or polyclonal. The solid support may then be washed
 5 with suitable buffers to give an immobilized gp120-labeled immunoglobulin-like molecule antibody complex. The label on the fusion protein may then be detected to give a measure of endogenous gp120 and, thereby, the presence of HIV.

This aspect of the invention relates to a method for detecting HIV or SIV viral infection in a sample comprising

10 (a) contacting a sample suspected of containing gp120 with a fusion protein or immunoglobulin-like molecule comprising CD4, or fragment thereof which binds to gp120, and the Fc portion of an immunoglobulin chain,

(b) detecting whether a complex is formed.

15 The invention also relates to a method of detecting gp120 in a sample, further comprising

(c) contacting the mixture obtained in step (a) with an Fc binding molecule, such as an antibody, protein A, or protein G, which is immobilized on a solid phase support and is specific for the hybrid fusion protein, to give a gp120 fusion protein-immobilized antibody complex

(d) washing the solid phase support obtained in step (c) to remove unbound fusion protein,

20 (e) and detecting the label on the hybrid fusion protein.

Of course, the specific concentrations of detectably labeled immunoglobulin-like molecule (or fusion protein) and gp120, the temperature and time of incubation, as well as other assay conditions may be varied, depending on various factors including the concentration of gp120 in the sample, the nature of the
 25 sample, and the like. Those skilled in the art will be able to determine operative and optimal assay conditions for each determination by employing routine experimentation.

Other such steps as washing, stirring, shaking, filtering and the like may be added to the assays as is customary or necessary for the particular situation.

One of the ways in which the immunoglobulin-like molecule or fusion protein of the present invention
 30 can be detectably labeled is by linking the same to an enzyme. This enzyme, in turn, when later exposed to its substrate, will react with the substrate in such a manner as to produce a chemical moiety which can be detected as, for example, by spectrophotometric, fluorometric or by visual means. Enzymes which can be used to detectably label the immunoglobulin-like molecule or fusion protein of the present invention include, but are not limited to, malate dehydrogenase, staphylococcal nuclease, delta-V-steroid isomerase, yeast
 35 alcohol dehydrogenase, alpha-glycerophosphate dehydrogenase, triose phosphate isomerase, horseradish peroxidase, alkaline phosphatase, asparaginase, glucose oxidase, beta-galactosidase, ribonuclease, urease, catalase, glucose-VI-phosphate dehydrogenase, glucoamylase and acetylcholine esterase.

The immunoglobulin-like molecule or fusion protein of the present invention may also be labeled with a radioactive isotope which can be determined by such means as the use of a gamma counter or a
 40 scintillation counter or by autoradiography. Isotopes which are particularly useful for the purpose of the present invention are: ^3H , ^{125}I , ^{131}I , ^{32}P , ^{35}S , ^{14}C , ^{51}Cr , ^{36}Cl , ^{57}Co , ^{58}Co , ^{59}Fe and ^{75}Se .

It is also possible to label the immunoglobulin-like molecule or fusion protein with a fluorescent compound. When the fluorescently labeled immunoglobulin-like molecule is exposed to light of the proper wave length, its presence can then be detected due to the fluorescence of the dye. Among the most
 45 commonly used fluorescent labelling compounds are fluorescein isothiocyanate, rhodamine, phycoerythrin, phycocyanin, allophycocyanin, o-phthaldehyde and fluorescamine.

The immunoglobulin-like molecule or fusion protein of the invention can also be detectably labeled using fluorescence emitting metals such as ^{152}Eu , or others of the lanthanide series. These metals can be attached to the immunoglobulin-like molecule or fusion protein using such metal chelating groups as
 50 diethylenetriaminepentaacetic acid (DTPA) or ethylenediaminetetraacetic acid (EDTA).

The immunoglobulin-like molecule or fusion protein of the present invention also can be detectably labeled by coupling it to a chemiluminescent compound. The presence of the chemiluminescent-tagged immunoglobulin-like molecule or fusion protein is then determined by detecting the presence of luminescence that arises during the course of a chemical reaction. Examples of particularly useful chemiluminescent labeling compounds are luminol, isoluminol, thermotropic acridinium ester, imidazole, acridinium salt and oxalate ester.

Likewise, a bioluminescent compound may be used to label the immunoglobulin-like molecule or fusion protein of the present invention. Bioluminescence is a type of chemiluminescence found in biological

systems in which a catalytic protein increases the efficiency of the chemiluminescent reaction. The presence of a bioluminescent protein is determined by detecting the presence of luminescence. Important bioluminescent compounds for purposes of labeling are luciferin, luciferase and aequorin.

Detection of the immunoglobulin-like molecule or fusion protein may be accomplished by a scintillation counter, for example, if the detectable label is a radioactive gamma emitter, or by a fluorometer, for example, if the label is a fluorescent material. In the case of an enzyme label, the detection can be accomplished by colorimetric methods which employ a substrate for the enzyme. Detection may also be accomplished by visual comparison of the extent of enzymatic reaction of a substrate in comparison with similarly prepared standards.

The assay of the present invention is ideally suited for the preparation of a kit. Such a kit may comprise a carrier means being compartmentalized to receive in close confinement therewith one or more container means such as vials, tubes and the like, each of said container means comprising the separate elements of the immunoassay. For example, there may be a container means containing a solid phase support, and further container means containing the detectably labeled immunoglobulin-like molecule or fusion protein in solution. Further container means may contain standard solutions comprising serial dilutions of analytes such as gp120 or fragments thereof to be detected. The standard solutions of these analytes may be used to prepare a standard curve with the concentration of gp120 plotted on the abscissa and the detection signal on the ordinate. The results obtained from a sample containing gp120 may be interpolated from such a plot to give the concentration of gp120.

The immunoglobulin-like molecule or fusion protein of the present invention can also be used as a stain for tissue sections. For example, a labeled immunoglobulin-like molecule comprising CD4 or fragment thereof which binds to gp120 may be contacted with a tissue section, e.g., a brain biopsy specimen. This section may then be washed and the label detected.

The following examples are illustrative, but not limiting the method and composition of the present invention. Other suitable modifications and adaptations which are obvious to this skill in the art are within the spirit and scope of this invention.

EXAMPLES

Example 1: Preparation of CD4-Ig cDNA Constructs

The extracellular portion of the CD4 molecule (See Madden, P.J., *et al.*, *Cell* 42:93-104 (1985)) was fused at three locations in a human IgG1 heavy chain constant region gene by means of a synthetic splice donor linker molecule. To exploit the splice donor linker, a BamHI linker having the sequence CGCGGATCCGCG was first inserted at amino acid residue 395 of the CD4 precursor sequence (nucleotide residue 1295). A synthetic splice donor sequence

GATCCCGAGGGTGAGTACTA
GGCTCCCACTCATGATTCGA

bounded by BamHI and HindIII complementary ends was created and fused to the HindIII site in the intron preceding the CH1 domain, to the EcoRI site in the intron preceding the hinge domain, and to the BamI site preceding the CH2 domain of the IgG1 genomic sequence. Assembly of the chimeric genes by ligation at the BamHI site afforded molecules in which either the variable (V) region, the V+CH1 regions, or the V, CH1 and hinge regions were replaced by CD4. In the last case, the chimeric molecule is expected to form a monomer structure, while in the former, a dimeric molecule is expected.

On such genetic construct which contains the DNA sequence which encodes CD4 linked to human IgG1 at the Hind3 site upstream of the CH1 region (fusion protein CD4H γ 1) is depicted in Table 1. The plasmid containing this genetic construct (pCD4H γ 1) has been deposited in *E. coli* (MC1061/P3) at the American Type Culture Collection (ATCC) under the terms of the Budapest Treaty and given accession number 67611.

A second genetic construct which contains the DNA sequence which encodes CD4 linked to human IgG1 at the Esp site upstream of the hinge region (fusion protein CD4E γ 1) is depicted in Table 2. The

plasmid containing this genetic construct (pCD4E γ 1) has been deposited in E. coli (MC1061/P3) at the ATCC under the terms of the Budapest Treaty and given accession number 67610.

A third genetic construct which contains the DNA sequence which encodes CD4 linked to human IgM at the Mst2 site upstream of the CH1 region (fusion protein CD4M μ) is depicted in Table 3. The plasmid
5 containing this genetic construct (PCD4M μ) has been deposited in E. coli (MC1061/P3) at the ATCC under the terms of the Budapest Treaty and given accession number 67608.

A fourth genetic construct which contains the DNA sequence which encodes CD4 linked to human IgM at the Pst site upstream of the CH2 region (fusion protein CD4P μ) is depicted in Table 4. The plasmid
10 containing this genetic construct (PCD4P μ) has been deposited in E. coli (MC1061/P3) at the ATCC under the terms of the Budapest Treaty and given accession number 67609.

A fifth genetic construct which contains the DNA sequence which encodes CD4 linked to human IgG1 at the Ban1 site downstream from the hinge region (fusion protein CD4B γ 1) is depicted in Table 5.

Two similar constructs were prepared from the human IgM heavy chain constant region by fusion with the introns upstream of the μ CH1 and CH2 domains at an MstII site and a PstI site respectively. The
15 fusions were made by joining the PstI site of the CD4/IgG1 construct fused at the Esp site in IgG1 gene to the MstII and Pst sites in the IgM gene. In the first instance, this was performed by treatment of the Pst end with T4 DNA Polymerase and the MstII end with E. coli DNA Polymerase, followed by ligation; and in the second instance, by ligation alone.

Immunoprecipitation of the fusion proteins with a panel of monoclonal antibodies directed against CD4
20 epitopes showed that all of the epitopes were preserved. A specific high affinity association is demonstrated between the chimeric molecules and HIV envelope proteins expressed on the surface of cells transfected with an attenuated (reverse transcriptase deleted) proviral construct.

25

30

35

40

45

50

55

10

5
10
15
20
25
30
35
40
45
50
55

5
 M M
 B B
 O O
 2 2
 H I N F 1
 241 CCCAGAAGAAGAGCATACAATTCCACTGGAAAACTCCAACCAGATAAAGATTCTGGGAA 300

 GGGTCTTCTTCTCGTATGTTAAGGTGACCTTTTTGAGGTGGTCTATTCTAAGACCCTT
 10
 Q K K S I Q F H W K N S N Q I K I L G N -
 B S S F H
 NBS AA A A N H I
 15 LAP O VU L U U H N
 AN1 K A9 U 3 D A F
 422 1 26 1 A 2 1 1
 20 301 ATCAGGGCTCCTTCTTAATAAGGTCCATCCAAGCTGAATGATCGCGCTGACTCAAGAA 360

 TAGTCCCGAGGAAGAATTGATTTCCAGGTAGGTTGACTTACTAGCGCGACTGAGTTCTT
 Q G S F L T K G P S K L N D R A D S R R -
 25 S S H H
 MANAS BA I A I D
 BVLUT CU N F N D
 QAA9Y L3 F L F E
 22461 1A 1 2 1 1
 30 361 GAAGCCTTTGGGACCAAGGAACTTCCCCCTGATCATCAAGAATCTTAAGATAGAAGACT 420

 CTTCGGAAACCCTGGTTCCTTTGAAGGGGGACTAGTAGTTCTTAGAATTCTATCTTCTGA
 S L W D Q G N F P L I I K N L K I E D S -
 35
 S
 M M AMAM M
 B N VNUN A
 40 O L AL9L E
 2 1 2161 1
 45 421 CAGATACTTACATCTGTGAAGTGAGGACCAGAAGGAGGTGCAATTGCTAGTGTTCG 480

 GTCTATGAATGTAGACACTTCACCTCCTGGTCTTCCTCCTCCACGTTAACGATCACAAGC
 D T Y I C E V E D Q K E E V Q L L V F G -
 50
 55

5
 B S
 S P
 M Y
 1 1
 481 GATTGACTGCCAACTCTGACACCCACCTGCTTCAGGGGCAGAGCCTGACCCTGACCTTGG 540
 CTAAGTACGGTTGAGACTGTGGGTGGACGAAGTCCCCGTCTCGGACTGGGACTGGAACC
 10 L T A N S D T H L L Q G Q S L T L T L E -
 15 B BS
 BS SC D M H
 AP TR D N I S
 N1 NF E L F Y
 22 11 1 1 1 1
 / /
 20 AGAGCCCCCTGGTAGTAGCCCCCTCAGTGCAATGTAGGAGTCCAAGGGGTAAAAACATAC 600
 541 TCTCGGGGGGACCATCATCGGGGAGTCACGTTACATCCTCAGGTTCCCCATTTTGTATG
 S P P G S S P S V Q C R S P R G K N I Q -
 25 M MD ASP A BBH S B BS
 B ND LPV L APTIAR T A L TR
 O LE UBU U N1NACF X N A NF
 2 11 122 1 221111 1 1 4 11
 // / /// /
 30 AGGGGGGGAAGACCCTCTCCGTGTCTCAGCTGGAGCTCCAGGATAGTGGCACCTGGACAT
 601 TCCCCCCTTCTGGGAGAGGCACAGAGTCGACCTCGAGGTCCTATCACCGTGGACCTGTA 660
 G G K T L S V S Q L E L Q D S G T W T C -
 40 N
 NS M NM A
 LP B HA L
 AH O EE U
 31 2 11 1
 /
 45 GCACTGTCTTGCAGAACCAGAAGAAGGTGGAGTTCAAAATAGACATCGTGGTGTAGCTT 720
 661 CGTGACAGAACGTCTTGGTCTTCTCCACCTCAAGTTTTATCTGTAGCACCACGATCGAA
 T V L Q N Q K K V E F K I D I V V L A F -
 50
 55

5
10
15
20
25
30
35
40
45
50
55

55

5
 F D M I A PS H
 O D A N V ADPA I
 K E E F A VRUJ N
 1 1 3 1 1 AAM9 F
 2216 1
 ///

10
 1201 AGGCGGGGATGTGGCAGTGTCTGCTGAGTGACTCGGGACAGGTCTGCTGGAATCCAACA 1260
 TCCGCCCCCTACACCGTCACAGACGACTCACTGAGCCCTGTCCAGGACGACCTTAGGTTGT
 A G M W Q C L L S D S G Q V L L E S N I -

15
 S SA BHF BS H
 ANA HNCP SGNMAANXA RSD I A
 VLU PCRA PIUNMULHV SCD N L
 AA9 AIFL 1ADLH3A0A AAE D U
 236 2111 21211A421 111 3 1
 20
 TCAAGGTTCTGCCACATGGTCCACCCCGGTGCACGCGGATCCCAGGGTGAGTACTAAG
 1261 AGTTCCAAGACGGGTGTACCAGGTGGGGCCACGTGCGCCTAGGGCTCCCACTCATGATTC 1320
 K V L P T W S T P V H A D P E

25
 BS B
 H SC HS S M M D S
 P A TR AT T N N D P
 H E NF EU Y L L E M
 1 3 11 31 1 1 1 1
 / / / / /

30
 CTTTCTGGGGCAGGCCAGGCCTGACCTTGCTTTGGGGCAGGGAGGGGGCTAAGGTGAGG
 1321 GAAAGACCCCGTCCGGTCCGGACTGGAACCGAAACCCCGTCCCTCCCCGATTCCACTCC 1380
 B A BH B P H
 BASHBHNN P SG N BS F H
 AHPHBAPAL A PI L AP L G
 40
 NAMEEHRA L 1A A N1 M A
 121112114 1 21 3 22 1 1
 / / / / /

45
 1381 CAGGTGGCGCCAGCAGGTGCACACCCAATGCCCATGAGCCCAGACACTGGACGCTGAACC 1440
 GTCCACCGCGGTCTGCCACGTGTGGGTTACGGGTACTCGGGTCTGTGACCTGCGACTTGG
 F M BS S B SS B S FN
 N DNHA H SMAAHNABSAC NS
 U N TR RLAU H TNUUALPAPLR UP
 D L NF AAE9 A NL99EAAN1UF DB
 50
 2 1 11 2436 1 11663412211 22
 / / / / /

55
 1441 TCGCGGACAGTTAAGAACCCAGGGGCTCTGCGCCTGGGGCCAGCTCTGTCCCACACCGC 1500
 AGCGCCTGTCAATTCTTGGGTCCCCGGAGACGCGGACCCGGGTGAGACAGGGTGTGGCG

MS BNN F BSS BS
 AA ALL NM S BMDMHNABSAA SCB
 EC NAA UN T BBRNALPAPUJ TRA
 32 134 4L Y VOALEAAN199 NFN
 / / H1 1 12213412266 111
 / / / / /
 1501 GGTACATGGCACCACCTCTCTTGCAGCCTCCACCAAGGGCCCATCGGTCTTCCCCCTGG
 ----- 1560
 CCAGTGTACCGTGGTGGAGAGAACGTCGGAGGTGGTTCCCGGGTAGCCAGAAGGGGGACC
 A S T K G P S V F P L A -

BH B NFS BS F BS
 M MSG MSB SNAH SC N SC
 N N NPI NPB PUUA TR U TR
 L L L1A L1V B49E NF 4 NF
 4 1 121 121 2H63 11 H 11
 / / / / /

1561 CACCCTCCTCCAAGAGCACCTCTGGGGGCACAGCGGCCCTGGGCTGCCTGGTCAAGGACT
 ----- 1620
 GTGGGAGGAGGTTCTCGTGGAGACCCCCGTGTCGCCGGGACCCGACGGACCAGTTCCTGA
 P S S K S T S G G T A A L G C L V K D Y -

H M T H D BANHBHN NF A BH
 P A T P D AHAHBAL SN P SG
 A E H H E NARAEAA B4 L 1A
 2 3 1 1 1 1211124 2H 1 21
 / / / / /

1621 ACTTCCCCGAACCGGTGACGGTGTCTGGAAGTCAAGCGCCCTGACCAGCGGCGTGCACA
 ----- 1680
 TGAAGGGGCTTGGCCACTGCCACAGCACCTTGAGTCCGCGGGACTGGTCCGCCACAGTGT
 F P E P V T V S W N S G A L T S G V H T -

S H F B B
 HNC DM I M D N M SM B
 PCR DS N N D U N TA B
 AIF ET F L E 4 L EE V
 211 12 1 1 1 H 1 23 1
 / / / / /

1681 CCTTCCCGGCTGTCTACAGTCTCAGGACTCTACTCCCTCAGCAGCGTGGTGACCGTGC
 ----- 1740
 GGAAGGGCCGACAGGATGTCAGGAGTCTGAGATGAGGGAGTCGTCGCACCACTGGCAGC
 F P A V L Q S S G L Y S L S S V V T V P -

B F B B H
 SH N ASM B NSB M I
 PP U LTN A LPB A N
 1H 4 UXL N A1V E F
 21 H 111 1 421 2 1
 / / / / /

1741 CCTCCAGCAGCTTGGGCACCCAGACCTACATCTGCAACGTGAATCACAAGCCCAGCAACA
 ----- 1800
 GGAGGTCGTGCAACCCGTGGGTCTGGATGTAGACGTTGCACTTAGTGTTCGGGTCTGTGT
 S S S L G T Q T Y I C N V N H K P S N T -

S M HM HM
 T N AN PN
 Y L EL HL
 1 1 31 11
 CCAAGGTGGACAAGAAAGTTGGTGAGAGGCCAGCACAGGGAGGGAGGGTGTCTGCTGGAA
 1801 ----- 1860
 GGTTCACCTGTTCTTTCAACCACTCTCCGGTCGTGTCCCTCCCTCCACAGACGACCTT
 K V D K K V
 E BS SS F BS F
 DE CHH F SC HHNCF N BSC N
 DS OHA O TR PGCRA U BTR U
 EP 4AE K NF AAIFN 4 VNF 4
 11 712 1 11 21111 H 111 H
 / / / /
 GCAGGCTCAGCGCTCCTGCCTGGACGCATCCCGGCTATGCAGCCCCAGTCCAGGGCAGCA
 1861 ----- 1920
 CGTCCGAGTCGCGAGGACGGACCTGCGTAGGGCCGATACGTGGGGTTCAGGTCCCGTCGT
 S S M MNDM
 DBHMHNA HMNCN N NLDB
 RBABPLU PNCRL L LAEO
 AVEOHA9 ALIFA 1 1312
 2132146 21114
 // //
 AGGCAGGCCCCGTCTGCCTCTTCACCCGGAGCCTCTGCCCGCCCCACTCATGCTCAGGGA
 1921 ----- 1980
 TCCGTCCGGGGCAGACGGAGAAGTGGCCCTCGGAGACGGGCGGGGTGAGTACGAGTCCCT
 BS P B BS
 SC F M B N S SC
 TR L A A L P TR
 NF M E N A 1 NF
 11 1 1 1 4 2 11
 /
 GAGGGTCTTCTGGCTTTTTCCCAGGCTCTGGGCAGGCACAGGCTAGGTGCCCTAACCCA
 1981 ----- 2040
 CTCCCAGAAGACCGAAAAAGGGTCCGAGACCCGTCCGTGTCCGATCCACGGGGATTGGGT
 S B B B S
 DHA S DBS S M HNC A
 RAU P DAP P N PCR V
 AE9 M EN1 M L AIF A
 236 1 122 1 1 211 2
 / / /
 GGCCCTGCACACAAAGGGGCAGGTGCTGGGCTCAGACCTGCCAAGAGCCATATCCGGGAG
 2041 ----- 2100
 CCGGGACGTGTGTTTCCCGTCCACGACCCGAGTCTGGACGGTTCTCGGTATAGGCCCTC

PS
 DNPA D H D A M
 5 RLUU D A D L N
 AAM9 E E U L
 2416 1 3 1 1 1
 / / /
 10 2101 GACCTGCCCCCTGACCTAAGCCCAACCCAAAGGCCAAACTCTCCACTCCCTCAGCTCGGA 2160
 CTGGGACGGGGACTGGATTGGGTGGGGTTTCCGGTTTGAGAGGTGAGGGAGTCGAGCCT

 H B
 15 I M MM P BS
 N N AB S AP
 F L EO T N1
 1 1 32 1 22
 / /
 20 2161 CACCTTCTCTCCTCCCAGATTCCAGTAACCTCCCAATCTTCTCTCTGCAGAGCCCAAATCT 2220
 GTGGAAGAGAGGAGGGTCTAAGGTCATTGAGGGTTAGAAGAGAGACGTCTCGGGTTTAGA

 E P K S -

 25 M N BBS BS M
 A NS SSC SC HS N
 E LP PTR TR AT
 3 AH 1NF NF EU L
 31 211 11 31 1
 / / /
 30 2221 TGTGACAAAACCTCACACATGCCACCGTGCCAGGTAAGCCAGCCCAGGCCTCGCCCTCC 2280
 ACACTGTTTTGAGTGTGTACGGGTGGCAGGGTCCATTGGTGGGTCCGGAGCGGGAGG
 C D K T H T C P P C P

 35 A M B N SM F BS S S S
 L N A L PA O SC F DHNA HNC
 U L N A 1E K TR A RALU PCR
 40 1 1 1 4 21 1 11 1 2346 211
 / / /
 2281 AGCTCAAGGCGGGACAGGTGCCCTAGAGTAGCCTGCATCCAGGGACAGGCCCCAGCCGGG 2340
 TCGAGTTCCGCCCTGTCCACGGGATCTCATCGGACGTAGGTCCCTGTCCGGGGTCGGCCC

 45 A M M M D M BS S
 F A B N D N SC M ANA M
 L E O L E L TR B VLU B
 50 3 2 2 1 1 1 11 2 246 2
 / /
 2341 TGCTGACACGTCCACCTCCATCTCTTCTCAGCACCTGAACTCCTGGGGGGACCGTCAGT 2400
 ACGACTGTGCAGGTGGAGGTAGAGAAGGAGTCGTGGACTTGAGGACCCCCCTGCCAGTCA

 55 A P E L L G G P S V -

5
 M S S SS
 N T AN M HMANNAC DM M
 L Y UL N PNVCLUR DS A
 1 1 3A L ALAIA9F ET E
 1 A3 1 2121461 12 3
 // // // //
 2401 CTCCTCTTCCCCCAAACCAAGGACACCTCATGATCTCCCGGACCCCTGAGGTCAC 2460
 10 GAAGGAGAAGGGGGGTTTTGGGTTCCTGTGGGAGTACTAGAGGGCCTGGGGACTCCAGTG
 F L F P P K P K D T L M I S R T P E V T -
 15 N M M DM M RM M
 NS LP A N DS B SA N
 AH E L ET O AE L
 31 2 1 12 2 12 1
 / /
 20 ATGCGTGGTGGTGGACGTGAGCCACGAAGACCCCTGAGGTCAAGTTCAACTGGTACGTGGA 2520
 2461 TACGCACCACCACCTGCACTCGGTGCTTCTGGGACTCCAGTTCAAGTTGACCATGCACCT
 C V V V D V S H E D P E V K F N W Y V D -
 25 F FN
 M N NSS R M R
 N U UPA S A S
 L 4 DBC A E A
 1 H 222 1 2 1
 //
 30 CGGCGTGGAGGTGCATAATGCCAAGACAAAGCCGCGGGAGGAGCAGTACAACAGCACGTA 2580
 2521 GCCGCACCTCCACGTATTACGGTTCTGTTTCGGCGCCCTCCTCGTCATGTTGTCGTGCAT
 G V E V H N A K T K P R E E Q Y N S T Y -
 35 S BS
 HNC HH M SC R
 PCR GP N TR S
 40 AIF AH L NF A
 211 11 1 11 1
 / /
 45 CCGGGTGGTCAGCGTCCTCACCCTCCTGCACCAGGACTGGCTGAATGGCAAGGAGTACAA 2640
 2581 GGCCCAACAGTCGCAGGAGTGGCAGGACGTGGTCCTGACCGACTTACCGTTCTCATGTT
 R V V S V L T V L H Q D W L N G K E Y K -
 50 M T
 N A
 L Q
 1 1
 2641 GTGCAAGGTCTCCAACAAAGCCCTCCAGCCCCCATCGAGAAAACCATCTCCAAGCCAA 2700
 55 CACGTTCCAGAGGTTGTTTCGGGAGGGTCCGGGGTAGCTCTTTTGGTAGAGGTTTCGGTT
 C K V S N K A L P A P I E K T I S K A K -

5
 P S
 ADNNPMA
 VRLLUNU
 AAAAML9
 2244116
 S
 A H M N
 U A N L
 9 E L A
 6 3 1 3
 HHN
 APA
 EAE
 321
 S
 BSAH
 GFUA
 LI9E
 1163
 // // /
 10 2701 AGGTGGGACCCGTGGGGTGGGAGGGCCACATGGACAGAGGCCGGCTCGGCCACCCCTCTG

 TCCACCCTGGGCACCCACGCTCCCGGTGTACCTGTCTCCGGCCGAGCCGGGTGGGAGAC 2760
 15
 D M M N S R M F N A B
 D N A P S N U V B
 E L E B A L 4 A V
 1 1 3 2 1 1 H 1 1
 20 2761 CCCTGAGAGTGACCGCTGTACCAACCTCTGTCTACAGGGCAGCCCCGAGAACCACAGGT

 GGGACTCTCACTGGCGACATGGTTGGAGACAGGATGTCCCGTCGGGGCTCTTGGTGTCCA 2820
 G Q P R E P Q V -
 25
 R F SS BS BS
 S O AHNNCCS SC SC
 A K VPCCRRM L O TR TR
 1 1 AAIFFA U K NF NF
 1211111 1 1 11 11
 30 2821 GTACACCCTGCCCCCATCCCGGGATGAGCTGACCAAGAACCAGGTCAGCCTGACCTGCCT

 CATGTGGGACGGGGGTAGGGCCCTACTCGACTGGTTCTTGGTCCAGTCGGACTGGACGGA 2880
 Y T L P P S R D E L T K N Q V S L T C L -
 35
 B F
 S N H
 P U P
 M 4 A
 1 H 2
 40 2861 GGTCAAAGGCTTCTATCCCAGCGACATCGCCGTGGAGTGGGAGAGCAATGGGCAGCCGGA

 CCAGTTTCCGAAGATAGGGTGGCTGTAGCGGCACCTCACCTCTCGTTACCCGTGCGCCT 2940
 V K G F Y P S D I A V E W E S N G Q P E -
 45
 H
 B M I M N H
 B N N B L P
 V L F O A H
 1 1 1 2 4 1
 50 2941 GAACAACTACAAGACCACGCCTCCCGTGGTGGACTCCGACGGCTCCTTCTTCTCTACAG

 CTTGTTGATGTTCTGGTGGGAGGGCAGACCTGAGGCTGCCGAGGAAGAAGGAGATGTC 3000
 55
 N N Y K T T P P V L D S D G S F F L Y S -

	M A	B	F	MBX	S	
5	N L	P	NM	ABM	NF	M
	L U	M	UB	EVN	LA	N
	1 1	1	40	211	AN	L
			H2		31	1
					/	
10	3001	CAAGCTCACCGTGGACAAGAGCAGGTGGCAGCAGGGGAACGTCTTCTCATGCTCCGTGAT				3060

		GTTTCGAGTGGCACCTGTTCTCGTCCACCGTCGTCCCCTTGCAGAAGAGTACGAGGGCACTA				
		K L T V D K S R W Q Q G N V F S C S V M -				
15					S	
	N N			M	M	HNC
	S L			B	N	PCR
	I A			O	L	AIF
20	1 3			2	1	211
					/	
	3061	GCATGAGGCTCTGCACAACCACTACACGCAGAAGAGCCTCTCCCTGTCTCCGGGTAAATG				3120

		CGTACTCCGAGACGTGTTGGTGATGTGCGTCTTCTCGGAGAGGGACAGAGGGCCATTAC				
25		H E A L H N H Y T Q K S L S L S P G K .				
		CXHHN				
		FMAPA				
30		RAEAE				
		13321				
		/				
	3121	AGTGCGACGGCCG				3133

35		TCACGCTGCCGGC				

Table 2

F N				S		B
N S	B	M	H	DHA		S
U P	B	N	G	RAU		T
4 B	V	L	A	AE9		X
H 2	1	1	1	236		1

GCCTGTTTGAGAAGCAGCGGGCAAGAAAGACGCAAGCCCAGAGGCCCTGCCATTTCTGTG
 1 ----- 60
 CGGACAAACTCTTCGTCGCCCGTTCTTTCTGCGTTCGGGTCTCCGGGACGGTAAAGACAC

B	PS		S		S	
DBS	ADNPA	D	DHNA	M	HM	HNC
DAP	VRUU	D	RALU	N	AN	PCR
EN1	AAAM9	E	AEA9	L	EL	AIF
122	22416	1	2346	1	31	211
/	/	/	/	/	/	/

GGCTCAGGTCCCTACTGGCTCAGGCCCTGCCTCCCTCGGCAAGGCCACAATGAACCGGG
 61 ----- 120
 CCGAGTCCAGGGATGACCGAGTCCGGGGACGGAGGGAGCCGTTCCGGTGTACTTGCCCC

				M	N	R	G	-
H			F		F			
I		B	N	HH	N	M	D	
N		B	U	HA	U	N	D	
F		V	4	AE	4	L	E	
1		1	H	12	H	1	1	

GAGTCCCTTTTAGGCACTTGCTTCTGGTGCTGCAACTGGCGCTCCTCCCAGCAGCCACTC
 121 ----- 180
 CTCAGGGAAAATCCGTGAACGAAGACCACGACGTTGACCGCGAGGAGGGTCTGTCGGTGAG

V P F R H L L L V L Q L A L L P A A T Q -

B	E	E			R	A
B	C	C			S	L
V	O	O			A	U
1	K	K			1	1

AGGGAAGAAAGTGGTGCTGGGCAAAAAAGGGGATACAGTGGAAGTACCTGTACAGCTT
 181 ----- 240
 TCCCTTTCTTTCACCACGACCCGTTTTTCCCTATGTCACCTTGACTGGACATGTCGAA

G K K V V L G K K G D T V E L T C T A S -

					H
	M	M			I
	B	B			N
	O	O			F
	2	2			1

CCCAGAAGAAGAGCATACAATTCCACTGGAAAACTCCAACCAGATAAAGATTCTGGGAA
 241 ----- 300
 GGGTCTTCTTCTCGTATGTTAAGGTGACCTTTTTGAGGTTGGTCTATTTCTAAGACCTT

Q K K S I Q F H W K N S N Q I K I L G N -

5
 M MD N BBH S B BS
 B ND ASP A BSSGSC S B N SC
 O LE UBU U N1NACF X N A NF
 2 11 122 1 221111 1 1 4 11
 // / /// /
 10
 601 AGGGGGGGAAGACCCTCTCCGTGTCTCAGCTGGAGCTCCAGGATAGTGGCACCTGGACAT 660
 TCCCCCCTTCTGGGAGAGGCACAGAGTCGACCTCGAGGTCTATCACCGTGGACCTGTA
 G G K T L S V S Q L E L Q D S G T W T C -
 15
 N
 NS M NM A
 LP B HA L
 AH O EE U
 31 2 11 1
 /
 20
 661 GCACCTGTCTTGCAAGACCAGAAGAAGGTGGAGTTCAAATAGACATCGTGGTGTCTAGCTT 720
 CGTGACAGAACGTCTTGGTCTTCTCCACCTCAAGTTTTATCTGTAGCACCACGATCGAA
 T V L Q N Q K K V E F K I D I V V L A F -
 25
 HS M M
 AT N N
 EU L L
 31 1 1
 /
 30
 721 TCCAGAAGGCCTCCAGCATAGTCTATAAGAAAGAGGGGGAACAGGTGGAGTTCTCCTTCC 780
 AGGTCTTCCGGAGGTCTGTATCAGATATTCTTTCTCCCCCTTGTCCACCTCAAGAGGAAGG
 Q K A S S I V Y K K E G E Q V E F S F P -
 35
 A A M
 L L N
 U U L
 1 1 1
 40
 781 CACTCGCCTTTACAGTTGAAAAGCTGACGGGCAGTGGCGAGCTGTGGTGGCAGGCGGAGA 840
 GTGAGCGGAAATGTCAACTTTTCGACTGCCCCTCACCCTCGACACCACCGTCCGCCTCT
 L A F T V E K L T G S G E L W W Q A E R -
 45
 P S
 H M FV A M
 P N LN U B
 H L ML 3 O
 1 1 11 A 2
 50
 841 GGGCTTCCTCCTCCAAGTCTTGGATCACCTTTGACCTGAAGAACAAGGAAGTGTCTGTAA 900
 CCCGAAGGAGGAGGTTTCAAGACCTAGTGAAACTGGACTTCTTGTTCTTCACAGACATT
 55
 A S S S K S W I T F D L K N K E V S V K -

B SM BS PS A H
 TA TRVRLUUD L P
 EE NFAAAM9E U H
 23 11224161 1 11

/ / / /
 AACGGGTTACCCAGGACCCCTAAGCTCCAGATGGGCAAGAAGCTCCCGCTCCACCTCACCC
 901 ----- 960
 TTGCCCAATGGGTCTCTGGGATTCTGAGGTCTACCCGTTCTTCGAGGGCGAGGTGGAGTGGG
 R V T Q D P K L Q M G K K L P L H L T L -

BS BSS
 M SC HS D M H SCAHM
 N TR AT D N P TRUAN
 L NF EU E L H NF9EL
 1 11 31 1 1 11631

/ /
 TGCCCCAGGCCTTGCCCTCAGTATGCTGGCTCTGGAAACCTCACCCCTGGCCCTTGAAGCGA
 961 ----- 1020
 ACGGGGTCCGGAACGGAGTCATACGACCGAGACCTTTGGAGTGGGACCGGGAACCTTCGCT
 P Q A L P Q Y A G S G N L T L A L E A K -

S BS
 F SC HD A
 A TR PD L
 N NF HE U
 1 11 1 1 1

/
 AAACAGGAAAGTTGCATCAGGAAGTGAACCTGGTGGTGATGAGAGCCACTCAGCTCCAGA
 1021 ----- 1080
 TTTGTCTTTCAACGTAGTCCTTCACTTGGACCACCCTACTCTCGGTGAGTCGAGGTCT
 T G K L H Q E V N L V V M R A T Q L Q K -

PS S
 M ADNPA DF AM DE A
 N VRLLUU DA LN DS L
 L AAAAM9 EN UL EP U
 1 224416 11 11 11 1

/ / / /
 AAAATTTGACCTGTGAGGTGTGGGGACCCACCTCCCTAAGCTGATGCTGAGCTTGAAC
 1081 ----- 1140
 TTTTAAACTGGACACTCCACACCCCTGGGTGGAGGGGATTGACTACGACTCGAAGCTTG
 N L T C E V W G P T S P K L M L S L K L -

M T H M DM
 N A P N DS
 L Q A L ET
 1 1 2 1 12

/
 TGGAGAACAAGGAGGCAAAGGTCTCGAAGCGGGGAGAAGCCGGTGTGGGTGCTGAACCTG
 1141 ----- 1200
 ACCTCTTGTTCTCCGTTTCCAGAGCTTCGCCCTCTTCGGCCACACCCACGACTTGGGAC
 E N K E A K V S K R E K P V W V L N P E -

5

	F	D	M	I	A		PS	H
	O	D	A	N	V		ADPA	I
	K	E	E	F	A		VRUU	N
	1	1	3	1	1		AAM9	F
							2216	1

10

1201 AGCGCGGGATGTGGCAGTGTCTGCTGAGTGACTCGGGACAGGTCCTGCTGGAATCCAACA 1260

TCCGCCCTACACCGTCACAGACGACTCACTGAGCCCTGTCCAGGACGACCTTAGGTTGT

A G M W Q C L L S D S G Q V L L E S N I -

15

	S	SA	BHF BS	H
	ANA	HNCP	SGNMAANXA	RSD I A
	VLU	PCRA	PIUNMULHV	SCD N L
	AA9	AIFL	1ADLH3A0A	AAE D U
	236	2111	21211A421	111 3 1

20

TCAAGGTTCTGCCCACATGGTCCACCCCGGTGCACGCGGATCCCCGAGGGTGAGTACTAAG

1261 AGTTC AAGACGGGTGTACCAGGTGGGGCCACGTGCGCCTAGGGCTCCCACTCATGATTC 1320

25

K V L P T W S T P V H A D P E

	E	BS	SS	F	BS	F
H	CHH	SC	HHNCF	N	BSC	N
P	OHA	TR	PGCRA	U	BTR	U
H	4AE	K NF	AAIFN	4	VNF	4
1	712	1 11	21111	H	111	H

30

CTTCAGCGCTCCTGCCTGGACGCATCCCGGCTATGCAGCCCCAGTCCAGGGCAGCAAGGC

1321 GAAGTCGCGAGGACGGACCTGCGTAGGGCCGATACGTGCGGGTTCAGGTCCCGTCGTTCCG 1380

35

S	S		
DBH/HNA	H/HNCN	M	MNDM
RB4SPLU	PNCRL	N	NLDB
AVEOHA9	A_IFA	L	LAEO
2132146	21114	1	1312

40

AGGCCCCGTCTGCCTCTTCACCCGGAGCCTCTGCCCCCCCCACTCATGCTCAGGGAGAGG

1381 TCCGGGGCAGACGGAGAAGTGGGCCTCGGAGACGGGCGGGGTGAGTACGAGTCCCTCTCC 1440

45

BS	P		B	BS	S
SC	F	M	B N	SCDHA	
TR	L	A	A L	TRRAU	
NF	M	E	N A	NFAE9	
11	1	1	1 4 2	11236	

50

GTCTTCTGGCTTTTTTCCAGGCTCTGGGCAGGCACAGGCTAGGTGCCCTAACCCAGGCC

1441 CAGAAGACCGAAAAAGGTCCGAGACCCGTCCGTGTCCGATCCACGGGGATTGGGTCCGG 1500

55

5
 B S P M 1
 B DBS DAP EN1 122 /
 B S M P N M L 1 1
 S HNC PCR AIF 211 /
 PS ADNPA VRLUUAAM9 22416 /
 10
 1501 CTGCACACAAAGGGGCAGGTGCTGGGCTCAGACCTGCCAAGAGCCATATCCGGGAGGACC 1560
 GACGTGTGTTTCCCCGTCCACGACCCGAGTCTGGACGGTTCTCGGTATAGGCCCTCCTGG
 15
 D D E 1
 H A E 3
 D A M D L N D U L 1 1 1
 1561 CTGCCCCCTGACCTAAGCCCCACCCCAAAGGCCAAACTCTCCACTCCCTCAGCTCGGACACC 1620
 GACGGGGACTGGATTGCGGTGGGGTTTCCGGTTTGAGAGGTGAGGGAGTCGAGCCTGTGG
 20
 H I M MM N N AB F L EO 1 1 32 /
 P B S BS M S AP M T N1 A 1 22 3 /
 1621 TTCTCTCCTCCCAGATTCCAGTAACTCCCAATCTTCTCTCTGCAGAGCCCAAATCTTGTG 1680
 AAGAGAGGAGGGTCTAAGGTCATTGAGGGTTAGAAGAGAGACGTCTCGGGTTTAGAACAC
 25
 30
 E P K S C D -
 35
 N NS LP AH 31 /
 BBS SSC PTR 1NF 211 /
 BS SC HS TR AT NF EU 11 31 /
 M A N L L U 1 1
 40
 1681 ACAAAAATCAGACATGCCCACTGTGCCCAAGGTAAGCCAGCCAGGCCTCGCCCTCCAGCT 1740
 TGTGTTTGAGTGTGTACGGGTGGCACGGGTCCATTGCGGTGCGGTCCGGAGCGGGAGGTCCA
 45
 K T H T C P P C P
 50
 M N L 1
 B N A N 1 4
 B SV PA 1E 21 1
 BS SC TR NF 11
 S F A N 1
 S DHNA RALU AEA9 2346
 S HNC PCR AIF 211
 1741 CAAGGCGGGACAGGTGCCCTAGAGTAGCCTGCATCCAGGGACAGGCCCCAGCCGGGTGCT 1800
 GTTCCGCCCTGTCCACGGGATCTCATCGGACGTAGGTCCCTGTCCGGGGTTCGGCCACGA
 55

5
 A M M M D M BS S
 F A B N D N SC M ANA M
 L E O L E L NF O AA9 O
 3 2 2 1 1 1 11 2 246 2
 1801 GACACGTCCACCTCCATCTCTTCCTCAGCACCTGAACTCCTGGGGGGACCGTCAGTCTTC 1860
 CTGTGCAGGTGGAGGTAGAGAAGGAGTCGTGGACTTGAGGACCCCCCTGGCAGTCAGAAG
 10
 A P E L L G G P S V F -
 15
 M S S N
 N T AN M HMANNAC DM M NS
 L Y UL N PNVCLUR DS A LP
 1 1 3A L ALAIA9F ET E AH
 1 A3 1 2121461 12 3 31
 1861 CTCTTCCCCCAAACCAAGGACACCCCTCATGATCTCCCGGACCCCTGAGGTACATGC 1920
 GAGAAGGGGGGTTTTGGGTTCTGTGGGAGTACTAGAGGGCCTGGGGACTCCAGTGTACG
 L F P P K P K D T L M I S R T P E V T C -
 25
 M M DM M RM M
 A N DS B SA N
 E L ET O AE L
 2 1 12 2 12 1
 1921 GTGGTGGTGGACGTGAGCCACGAAGACCCTGAGGTCAAGTTCAACTGGTACGTGGACGGC 1980
 CACCACCACCTGCACTCGGTGCTTCTGGGACTCCAGTTCAAGTTGACCATGCACCTGCCG
 V V V D V S H E D P E V K F N W Y V D G -
 35
 F FN S
 M N NES R M R HNC
 N U UPA S A S PCR
 L 4 DBC A E A AIF
 40 1 H 222 1 2 1 211
 1981 GTGGAGGTGCATAATGCCAAGACAAAGCCGCGGGAGGAGCAGTACAACAGCACGTACCGG 2040
 CACCTCCACGTATTACGGTTCTGTTTCGGCGCCCTCCTCGTCATGTTGTCGTGCATGGCC
 45
 V E V H N A K T K P R E E Q Y N S T Y R -
 50
 55

33

Table 3

5		F N				S		B
		N S	B	M	H	DHA		S
		U P	B	N	G	RAU		T
		4 B	V	L	A	AE9		X
		H 2	1	1	1	236		1
10						/		
	1	GCCTGTTTGAGAAGCAGCGGGCAAGAAAGACGCAAGCCCAGAGGCCCTGCCATTTCTGTG						
		CGGACAAACTCTTCGTCGCCCCGTTCTTTCTGCGTTCGGGTCTCCGGGACGGTAAAGACAC						60
15		B PS		S				S
		DBS ADNPA	D	DHNA	M	HM		HNC
		DAP VRLUU	D	RALU	N	AN		PCR
		EN1 AAAM9	E	AEA9	L	EL		AIF
		122 22416	1	2346	1	31		211
		/ / //		/		/		/
20		GGCTCAGGTCCCTACTGGCTCAGGCCCTGCCTCCCTCGGCAAGGCCACAATGAACCGGG						
	61	CCGAGTCCAGGGATGACCGAGTCCGGGGACGGAGGGAGCCGTTCCGGTGTTACTTGGCCC						120
25							M N R G -	
		H		F		F		
		I	B	N	HH	N M	D	
		N	B	U	HA	U N	D	
		F	V	4	AE	4 L	E	
30		1	1	H	12	H 1	1	
		GAGTCCCTTTTAGGCACTTGCTTCTGGTGCTGCAACTGGCGCTCCTCCCAGCAGCCACTC						
	121	CTCAGGGGAAAATCCGTGAACGAAGACCACGACGTTGACCGCGAGGAGGGTCGTCGGTGAG						180
35		V P F R H L L L V L Q L A L L P A A T Q -						
		B E E				R A		
		B C C				S L		
		V O O				A U		
40		1 K K				1 1		
		AGGGAAAGAAAGTGCTGCTGGGCAAAAAAGGGGATACAGTGGAAGTACCTGTACAGCTT						
	181	TCCCTTTCTTTCAACACGACCCGTTTTTTCCCTATGTCACCTTGACTGGACATGTGCGAA						240
45		G K K V V L G K K G D T V E L T C T A S -						
50								
55								

5
 241
 10
 15
 20
 25
 30
 361
 35
 40
 421
 45
 50
 55

M M
 B B
 C C
 2 2

H
 I
 N
 F
 1

CCCAGAAGAAGAGCATACAATTCCACTGCAAAACTCCAACCAGATAAAGATTCTGGGAA
 GGGTCTTCTTCTCGTATGTTAAGGTGACCTTTTGGAGGTGGTCTATTTCTAAGACCCTT

Q K K S I Q F H W K N S N Q I K I L G N -

B S
 NBS F AA A A F H I
 LAP O VU L U UH N
 AN1 K A9 U 3 D A F
 422 1 26 1 A 2 1 1

ATCAGGGCTCCTTCTTAATAAGGTCCATCCAAGCTGAATGATCGCGCTGACTCAAGAA
 TAGTCCCGAGGAAGAATTGATTTCCAGGTAGGTTGACTTACTAGCGCGACTGAGTTCTT

Q G S F L T K G P S K L N D R A D S R R -

S S H H
 MANAS BA I A I D
 BVLUT CU N F N D
 OAA9Y L3 F L F E
 22461 1A 1 2 1 1

GAAGCCTTTGGGACCAAGGAACTTCCCCCTGATCATCAAGAATCTTAAGATAGAAGACT
 CTTCGGAAACCCTGGTTCCTTTGAAGGGGGACTAGTAGTTCTTAGAATTCTATCTTCTGA

S L W D Q G N F P L I I K N L K I E D S -

S
 M M AMAM M
 B N VNUN A
 O L AL9L E
 2 1 2161 1

CAGATACTTACATCTGTGAAGTGGAGGACCAGAAGGAGGAGGTGCAATTGCTAGTGTTCTG
 GTCTATGAATGTAGACACTTCACCTCCTGGTCTTCTCCTCCACGTTAACGATCACAAGC

D T Y I C E V E D Q K E E V Q L L V F G -

5
 10
 15
 20
 25
 30
 35
 40
 45
 50
 55

B
S
P
M
1

S
T
Y
1

481 ----- 540

GATTGACTGCCAACTCTGACACCCACCTGCTTCAGGGGCAGAGCCTGACCCTGACCTTGG
 CTAAGTACGCGTTGAGACTGTGGGTGGACGAAGTCCCCGTCTCGGACTGGGACTGGAACC

L T A N S D T H L L Q G Q S L T L T L E -

B BS
 BS SC
 AP TR
 N1 NF
 22 11

D
 D
 E
 1

N
 N
 L
 1

H
 I
 N
 F
 1

S
 T
 Y
 1

541 ----- 600

AGAGCCCCCTGGTAGTAGCCCCCTCAGTGCAATGTAGGAGTCCAAGGGGTAAAAACATAC
 TCTCGGGGGGACCATCATCGGGGAGTCACGTTACATCCTCAGGTTCCCCATTTTGTATG

S P P G S S P S V Q C R S P R G K N I Q -

M MD ASP A BSSGSC S B N SC
 B ND LPV L APTIAR T A L TR
 O LE UBU U N1NACF X N A NF
 2 11 122 1 221111 1 1 4 11

// / /// /

601 ----- 660

AGGGGGGAAGACCCTCTCCGTGTCTCAGCTGGAGCTCCAGGATAGTGGCACCTGGACAT
 TCCCCCCTTCTGGGAGAGGCACAGAGTCGACCTCGAGGTCCTATCACCGTGGACCTGTA

G G K T L S V S Q L E L Q D S G T W T C -

N
 NS
 LP
 AH
 31

M
 B
 O
 2

NM A
 HA L
 EE U
 11 1

661 ----- 720

GCACTGTCTTGCAGAACCAGAAGAAGGTGGAGTTCAAAATAGACATCGTGGTGTAGCTT
 CGTGACAGAACGTCTTGGTCTTCTCCACCTCAAGTTTTATCTGTAGCACCACGATCGAA

T V L Q N Q K K V E F K I D I V V L A F -

HS M M
 AT N N
 EU L L
 31 1 1
 /
 5 TCCAGAAGGCCTCCAGCATAGTCTATAAGAAAGAGGGGGAACAGGTGGAGTTCTCCTTCC
 721 ----- 780
 AGGTCTTCCGGAGGTCGTATCAGATATTCTTCTCCCCCTTGTCACCTCAAGAGGAAGG
 10 Q K A S S I V Y K K E G E Q V E F S F P -
 A A M
 L L N
 U U L
 1 1 1
 15 CACTCGCCTTTACAGTTGAAAAGCTGACGGGCAGTGGCGAGCTGTGGTGGCAGGCGGAGA
 781 ----- 840
 GTGAGCGGAAATGTCAACTTTTCGACTGCCCGTCACCGCTCGACACCACCGTCCGCCTCT
 20 L A F T V E K L T G S G E L W W Q A E R -
 P S
 H M F V A
 P N L N U
 25 H L M L 3
 1 1 1 1 A
 841 ----- 900
 GGGCTTCCTCCTCCAAGTCTTGGATCACCTTTGACCTGAAGAACAAGGAAGTGTCTGTAA
 CCCGAAGGAGGAGGTTTCAAGCTAGTGGAAGCTGGACTTCTTGTTCCTTCACAGACATT
 30 A S S S K S W I T F D L K N K E V S V K -
 B BS PS
 SM SCADNPAD A A H
 35 TA TRVRLUUD L L P
 EE NFAAAV9E U U H
 23 11224161 1 1 1
 / / / /
 40 AACGGGTTACCCAGGACCCTAAGCTCCAGATGGGCAAGAAGCTCCCGCTCCACCTCACCC
 901 ----- 960
 TTGCCCCAATGGGTCCTGGGATTGAGGTCTACCCGTTCTTCGAGGGCGAGGTGGAGTGGG
 R V T Q D P K L Q M G K K L P L H L T L -
 BS BSS
 45 M SC HS D M H
 N TR AT D N P
 L NF EU E L H
 1 11 31 1 1 1
 / / / /
 50 TGCCCCAGGCCTTGCCTCAGTATGCTGGCTCTGGAAACCTCACCTGGCCCTTGAAGCGA
 961 ----- 1020
 ACGGGGTCCGGAACGGAGTCATACGACCGAGACCTTTGGAGTGGGACCGGGAACCTTCGCT
 55 P Q A L P Q Y A G S G N L T L A L E A K -

5
S BS
F SC
A TR
N NF
1 11
/

10
1021 AAACAGGAAAGTTGCATCAGGAAGTGAACCTGGTGGTGATGAGAGCCACTCAGCTCCAGA 1080
TTTGTCCCTTTCAACGTAGTCCTTCACTTGGACCACCACTACTCTCGGTGAGTCGAGGTCT
T G K L H Q E V N L V V M R A T Q L Q K -

15
PS S
ADNNPA DF AN DE A
VRLLUO DA LN DS L
AAAAV9 EN UL EP U
224416 11 11 11 1
///// / / /

20
1081 AAAATTTGACCTGTGAGGTGTGGGGACCCACCTCCCTAAGCTGATGCTGAGCTTGAAAC 1140
TTTTAAACTGGACACTCCACACCCCTGGGTGGAGGGGATTGCGACTACGACTCGAACTTTG
N L T C E V W G P T S P K L M L S L K L -

25
M T H M DW
N A P N DS
L Q A L ET
1 1 2 1 12
/

30
1141 TGGAGAACAAGGAGGCCAAAGGTCTCGAAGCGGGAGAAGCCGGTGTGGGTGCTGAACCCCTG 1200
ACCTCTTGTTCCCTCCGTTTCCAGAGCTTCGCCCTCTTCGGCCACACCCACGACTTGGGAC
E N K E A K V S K R E K P V W V L N P E -

35
H PS H
F D M I A ADPA I
O D A N V VRUU N
K E E F A AA9 F
1 1 3 1 1 2216 1
///

40
45 1201 AGGCGGGGATGTGGCAGTGTCTGCTGAGTGACTCGGGACAGGTCTGCTGGAATCCAACA 1260
TCCGCCCCCTACACCGTCACAGACGACTCACTGAGCCCTGTCCAGGACGACCTTAGGTTGT
A G M W Q C L L S D S G Q V L L E S N I -

50
55

5
S SA BHF BS H
ANA HNCP SGNMAANXA RSD I A
VLU PCRA PIUNMULHV SCD N L
AA9 AIFL 1ADLH3A0A AAE D U
236 2111 21211A421 111 3 1
// // // // //
TCAAGGTTCTGCCCACATGGTCCACCCCGGTGCACGCGGATCCCGAGGGTGAGTACTAAG
1261 ----- 1320
10 AGTTCCAAGACGGGTGTACCAGGTGGGGCCACGTGCGCCTAGGGCTCCCACTCATGATTC
K V L P T W S T P V H A D P E
15
E BS SS F BS F
H CHH F SC HHNCF N BSC N
P OHA O TR PGCRA U BTR U
H 4AE K NF AAIFN 4 VNF 4
1 712 1 11 21111 H 111 H
/ / // //
20 CTTCAGCGCTCCTGCCTGGACGCATCCCGGCTATGCAGCCCCAGTCCAGGGCAGCAAGGC
1321 ----- 1380
GAAGTCGCGAGGACGGACCTGCGTAGGGCCGATACGTCGGGGTCAGGTCCCGTCGTTCCG
25
S S W WNDV
DBHMHNA HMNCA N NLDV
RBABPLU PNCR L LAED
AVEOHA9 ALIFA L LAED
2132146 21114 1 1312
// // //
30 AGGCCCCGTCTGCCTCTTCACCCGGAGCCTCTGCCCCCCCCACTCATGCTCAGGGAGAGG
1381 ----- 1440
TCCGGGGCAGACGGAGAAGTGGCCCTCGGAGACGGGCGGGGTGAGTACGAGTCCCTCTCC
35
BS P B BS S
SC F M B N S SCDHA
TR L A A L P TRRAU
NF M E N A 1 NFAE9
11 1 1 1 4 2 11236
/ / //
40 GTCTTCTGGCTTTTTCCCAGGCTCTGGGCAGGCACAGGCTAGGTGCCCCTAACCCAGGCC
1441 ----- 1500
CAGAAGACCGAAAAAGGTCCGAGACCCGTCGTCGATCCACGGGGATTGGGTCCGG
45
B B B S PS
S DBS S M HNC ADNPA
P DAP P N PCR VRLUJ
M EN1 M L AIF AAAM9
1 122 1 1 211 22416
/ / //
50 CTGCACACAAAGGGGCAGGTGCTGGGCTCAGACCTGCCAAGAGCCATATCCGGGAGGACC
1501 ----- 1560
GACGTGTGTTTCCCGTCCACGACCCGAGTCTGGACGGTTCTCGGTATAGGCCCTCTCGG

5
D D E 1 H A E 3 D D E 1 A L U 1 M N L 1
1561 CTGCCCCGACCTAAGCCCCACCCCAAAGGCCAAACTCTCCACTCCCTCAGCTCGGACACC 1620
GACGGGGACTGGATTGCGGTGGGGTTTCCGGTTTGAGAGGTGAGGGAGTCGAGCCTGTGG
10 L P L T * A H P K G Q T L H S L S S D T -
C P * P K P T P K A K L S T P S A R T P -
A P D L S P P Q R P N S P L P Q L G H L -
15 H I M MM N N AB DF DO EK S F A N 1
1 1 32 11
20 TTCTCTCCTCCCAGATTCCAGTAACTCCCAATCTTCTCTCTCAGGGAGTGCATCCGCCCC 1680
1621 AAGAGAGGAGGGTCTAAGGTCATTGAGGGTTAGAAGAGAGAGTCCCTCACGTAGCGGGG
G S A S A P -
25
30 W N L 1 E C O R 1
1681 AACCTTTTCCCCCTCGTCTCCTGTGAGAATTCC.... 1714
TTGGGAAAAGGGGGAGCAGAGGACACTCTTAAGG....
35 T L F P L V S C E N S
40
45
50
55

5
 M M
 B B
 O O
 2 2
 H I N F 1
 241 CCCAGAAGAAGAGCATACAATTCCACTGGAAAACTCCAACCAJATAAAGATTCTGGGAA 300
 GGGTCTTCTTCTCGTATGTTAAGGTGACCTTTTTGAGGTTGGTCTATTTCTAAGACCCTT
 10 Q K K S I Q F H W K N S N Q I K I L G N -
 B S
 NBS F AA A S F H
 LAP O VU L U N H I
 15 AN1 K A9 U 3 D A F
 422 1 26 1 A 2 1 1
 / /
 301 ATCAGGGCTCCTTCTTAATAAGGTCCATCCAAGCTGAATGATCGCGCTGACTCAAGAA 360
 TAGTCCCGAGGAAGAATTGATTTCCAGGTAGGTTGACTTACTAGCGCGACTGAGTTCTT
 20 Q G S F L T K G P S K L N D R A D S R R -
 S S H H
 25 MANAS BA I A I D
 BVLUT CU N F N D
 OAA9Y L3 F L F E
 22461 1A 1 2 1 1
 / /
 30 GAAGCCTTTGGGACCAAGGAAACTTCCCCCTGATCATCAAGAATCTTAAGATAGAAGACT
 361 CTTGCGAAACCCTGGTTCCTTTGAAGGGGGACTAGTAGTCTTAGAATTCTATCTTCTGA 420
 S L W D Q G N F P L I I K N L K I E D S -
 S
 M M AMAM M
 B N VNUN A
 40 O L AL9L E
 2 1 2161 1
 //
 421 CAGATACTTACATCTGTGAAGTGGAGGACCAGAAGGAGGAGGTGCAATTGCTAGTGTTGG 480
 GTCTATGAATGTAGACACTTCACCTCCTGGTCTTCCTCCTCCAGTTAACGATCACAAGC
 45 D T Y I C E V E D Q K E E V Q L L V F G -
 50
 55

5
 B
S
P
M
1
S
T
Y
1
 481 GATTGACTGCCAACTCTGACACCCACCTGCTTCAGGGGCAGAGCCTGACCCTGACCTTGG 540
 10 CTAAGTACGGTTGAGACTGTGGGTGGACGAAGTCCCCGTCTCGGACTGGGACTGGAACC
 L T A N S D T H L L Q G Q S L T L T L E -
 15 B BS D M H S
 BS SC D M I S
 AP TR D N N T
 N1 NF E L F Y
 22 11 1 1 1 1
 / /
 20 AGAGCCCCCTGGTAGTACCCCTCAGTGCAATGTAGGAGTCCAAGGGGTAAAAACATAC 600
 541 TCTCGGGGGGACCATCATCGGGGAGTCACGTTACATCCTCAGGTTCCCCATTTTGTATG 600
 S P P G S S P S V Q C R S P R G K N I Q -
 25 M MD ASP A BSH S B BS
 B ND LPV L APTIAR T A L TR
 O LE UBU U NINACF X N A NF
 2 11 122 1 221111 1 1 4 11
 // / /// /
 30 AGGGGGGGAAGACCCTCTCCGTGTCTCAGCTGGAGCTCCAGGATAGTGGCACCTGGACAT 660
 601 TCCCCCCTTCTGGGAGAGGCACAGAGTCGACCTCGAGGTCCTATCACCCTGGACCTGTA 660
 G G K T L S V S Q L E L Q D S G T W T C -
 35 N
 NS M NW A
 LP B HA L
 AH O EE U
 40 31 2 11 1
 /
 45 GCACCTGTCTTGCAAGACCAGAAGAAGGTGGAGTTCAAAATAGACATCGTGGTGCTAGCTT 720
 661 CGTGACAGAACGTCTTGGTCTTCTTCCACCTCAAGTTTTATCTGTAGCACCACGATCGAA 720
 T V L Q N Q K K V E F K I D I V V L A F -
 50 HS M M
 AT N N
 EU L L
 31 1 1
 /
 55 TCCAGAAGGCCTCCAGCATAGTCTATAAGAAAGAGGGGGAACAGGTGCAGTTCTCCTTCC 780
 721 AGGTCTTCCGGAGGTCGTATCAGATATTCTTTCTCCCCCTTGTCCACCTCAAGAGGAAG 780
 Q K A S S I V Y K K E G E Q V E F S F P -

5
 A L M
 L U L
 1 1 1
 CACTCGCCTTTACAGTTGAAAAGCTGACGGGCAAGTGCCGAGCTGTGGTGGCAGGCGGAGA
 781 ----- 840
 GTGAGCGGAAATGTCAACTTTTCGACTGCCCCGTCACCGCTCGACACCACCGTCCGCCTCT
 10 L A F T V E K L T G S G E L W W Q A E R -

 P S
 H M F M A M
 P N L N U B
 15 H L M L 3 O
 1 1 1 1 A 2
 GGGCTTCCTCCTCCAAGTCTTGGATCACCTTTGACCTGAAGAACAAGGAAGTGTCTGTAA
 841 ----- 900
 CCCGAAGGAGGAGGTTTCAACCTAGTGAAACTGGACTTCTTGTTCCTTCACAGACATT
 20 A S S S K S W I T F D L K N K E V S V K -

 B BS PS
 SM SCADNPAD A A H
 25 TA TRVRLUUD L L P
 EE NFAAAM9E U U H
 23 11224161 1 1 1
 / / / /
 AACGGGTTACCCAGGACCCCTAAGCTCCAGATGGGCAAGAAGCTCCCGCTCCACCTCACCC
 901 ----- 960
 TTGCCCAATGGGTCCTGGGATTGAGGTCTACCCGTTCTTCGAGGGCGAGGTGGAGTGGG
 30 R V T Q D P K L Q M G K K L P L H L T L -

 BS BSS
 M SC HS D M H SCAHM
 35 N TR AT D N P TRUAN
 L NF EU E L H NF9EL
 1 11 31 1 1 1 11631
 / / / /
 TGCCCCAGGCCTTGCCCTCAGTATGCTGGCTCTGGAAACCTCACCTGGCCCTTGAAGCGA
 40 961 ----- 1020
 ACGGGGTCCGGAACGGAGTCATACGACCGAGACCTTTGGAGTGGGACCGGGAACCTTCGCT
 P Q A L P Q Y A G S G N L T L A L E A K -

 S BS
 F SC
 45 A TR
 N NF
 1 11
 /
 50 AAACAGGAAAGTTGCATCAGGAAGTGAACCTGGTGGTGTGATGAGAGCCACTCAGCTCCAGA
 1021 ----- 1080
 TTTGTCCTTTCAACGTAGTCCTTCACTTGGACCACCACTACTCTCGGTGAGTGGAGGTCT
 T G K L H Q E V N L V V M R A T Q L Q K -
 55

5 M PS S DF AM DE A
 N ADNNPA DA LN DS L
 L VRLLUU EN UL EP U
 1 224416 11 11 11 1
 // // /
 10 1081 AAAATTTGACCTGTGAGGTGTGGGGACCCACCTCCCTAAGCTGATGCTGAGCTTGAAC 1140
 TTTTAAACTGGACACTCCACACCCCTGGGTGGAGGGGATTGACTACGACTCGAACTTTG
 N L T C E V W G P T S P K L M L S L K L -
 15 M T H M DM
 N A P N DS
 L Q A L ET
 1 1 2 1 12
 /
 20 1141 TGGAGAACAAGGAGGCAAAGGTCTCGAAGCGGGAGAAGCCGGTGTGGGTGCTGAACCTG 1200
 ACCTCTTGTTCTCCGTTTCCAGAGCTTCGCCCTCTTCGGCCACACCCACGACTTGGGAC
 E N K E A K V S K R E K P V W V L N P E -
 25 F D M I A PS H
 O D A N V ADPA I
 K E E F A VRUU N
 1 1 3 1 1 AAV9 F
 2216 1
 ///
 30 1201 AGGCGGGGATGTGGCAGTGTCTGCTGAGTGACTCGGGACAGGTCTGCTGGAATCCAACA 1260
 TCCGCCCTACACCGTCACAGACGACTCACTGAGCCCTGTCCAGGACGACCTTAGGTTGT
 A G M W Q C L L S D S G Q V L L E S N I -
 35 S SA BHF BS H
 ANA HNCP SGNVAANXA RSD I A
 VLU PCRA PIUNMULHV SCD N L
 AA9 AIFL 1ADLH3A0A AAE D U
 236 2111 21211A421 111 3 1
 40 // // // // /
 1261 TCAAGGTTCTGCCCACATGGTCCACCCCGGTGCACGCGGATCCCAGGGTGAGTACTAAG 1320
 AGTTCCAAGACGGGTGTACCAGGTGGGGCCACGTGCGCCTAGGGCTCCCACTCATGATTC
 45 K V L P T W S T P V H A D P E
 50 E BS SS F BS F
 H CHH F SC HHNCF N BSC N
 P OHA O TR PGCRA U BTR U
 H 4AE K NF AAIFN 4 VNF 4
 1 712 1 11 21111 H 111 H
 / / // //
 1321 CTTCAAGGCTCCTGCCTGGACGCATCCCGGTATGCAGCCCCAGTCCAGGGCAGCAAGGC 1380
 55 GAAGTCGCGAGGACGGACCTGCGTAGGGCCGATACGTGGGGTCAGGTCCCGTCGTTCCG

5 S S M MNDV
DBHIAHNA HMNCN N N_DS
RBABPLU PNCRL N N_DS
AVEOHA9 ALIFA L LAEG
2132146 21114 1 1312
// // //

10 AGGCCCCGTCTGCCTCTTCACCCGGAGCCTCTGCCCCCCCCACTCATGCTCAGGGAGAGG
----- 1440
1381 TCCGGGGCAGACGGAGAAGTGGGCCTCGGAGACGGGCGGGGTGAGTACGAGTCCCTCTCC

15 BS P B BS S
SC F M B N S SCDHA
TR L A A L P TRRAU
NF M E N A 1 NFAE9
11 1 1 1 4 2 11236
/ /

20 GTCTTCTGGCTTTTTCCAGGCTCTGGGCAGGCACAGGCTAGGTGCCCTAACCCAGGCC
----- 1500
1441 CAGAAGACCGAAAAAGGGTCCGAGACCCGTCCGTGCCGATCCACGGGGATTGGGTCCGG

25 B B B S PS
S DBS S M HNC ADNPA
P DAP P N PCR VRLUU
M EN1 M L AIF AAAY9
1 122 1 1 211 22416
/ / /

30 CTGCACACAAAGGGGCAGGTGCTGGGCTCAGACCTGCCAAGAGCCATATCCGGGAGGACC
----- 1560
1501 GACGTGTGTTTCCCCGTCCACGACCCGAGTCTGGACGGTTCTCGGTATAGGCCCTCCTGG

35 D H D A M
D A D L N
E E E U L
1 3 1 1 1

40 CTGCCCCTGACCTAAGCCCCACCCAAAGGCCAAACTCTCCACTCCCTCAGCTCGGACACC
----- 1620
1561 GACGGGGACTGGATTGCGGTGGGGTTTCCGGTTTGAGAGGTGAGGGAGTGGAGCCTGTGG

45 H I M MM BP DE F
I N N AB BS DS AN
F L EO VT EP U4
1 1 32 11 11 1H
/ / /

50 TTCTCTCTCCAGATTCCAGTAACTCCCAATCTTCTCTGCACTGATTGCTGAGCTGC
----- 1680
1621 AAGAGAGGAGGGTCTAAGGTCATTGAGGGTTAGAAGAGAGACGTCACTAACGACTCGACG

V I A E L P -

50

Table 5

5		F N				S		B
		N S	B	M	H	DHA		S
		U P	B	N	G	RAU		T
		4 B	V	L	A	AE9		X
		H 2	1	1	1	236		1
10						/		
	1	GCCTGTTTGAGAAGCAGCGGGCAAGAAAGACGCAAGCCCAGAGGCCCTGCCATTTCTGTG						
		-----+-----+-----+-----+-----+-----+-----+ 60						
		CGGACAAACTCTTCGTCGCCCCGTTCTTTCTGCGTTCGGGTCTCCGGGACGGTAAAGACAC						
15		B PS		S				S
		DBS ADNPA	D	DHNA	M	HM		HNC
		DAP VRLUU	D	RALU	N	AN		PCR
		EN1 AAAM9	E	AEA9	L	EL		AIF
		122 22416	1	2346	1	31		211
		/ / //		/		/		/
20		GGCTCAGGTCCCTACTGGCTCAGGCCCTGCCTCCCTCGGCAAGGCCACAATGAACCGGG						
	61	-----+-----+-----+-----+-----+-----+-----+ 120						
		CCGAGTCCAGGGATGACCGAGTCCGGGGACGGAGGGAGCCGTTCCGGTGTTACTTGGCCC						
25						M N R G -		
		H		F		F		
		I		N		N M		D
		N	B	U	HH	U N		D
		F	V	4	HA	4 L		E
		1	1	H	AE	H 1		1
					12			
30		GAGTCCCTTTTAGGCACTTGCTTCTGGTGCTGCAACTGGCGCTCCTCCCAGCAGCCACTC						
	121	-----+-----+-----+-----+-----+-----+-----+ 180						
		CTCAGGGAAAATCCGTGAACGAAGACCACGACGTTGACCGCGAGGAGGGTCGTGGTGAG						
35		V P F R H L L L V L Q L A L L P A A T Q -						
		B E E				R A		
		B C C				S L		
		V O O				A U		
		1 K K				1 1		
40		AGGGAAAGAAAGTGGTGCTGGGCAAAAAAGGGGATACAGTGGAAGTACCTGTACAGCTT						
	181	-----+-----+-----+-----+-----+-----+-----+ 240						
		TCCCTTTCTTTCACCACGACCCGTTTTTCCCTATGTCACCTTGACTGGACATGTCGAA						
45		G K K V V L G K K G D T V E L T C T A S -						
			M M			H		
			B B			I		
			O O			N		
			2 2			F		
50						1		
		CCCAGAAGAAGAGCATACAATCCACTGGAAAACTCCAACCAGATAAAGATTCTGGGAA						
	241	-----+-----+-----+-----+-----+-----+-----+ 300						
		GGGTCTTCTTCTCGTATGTTAAGGTGACCTTTTTGAGGTGGTCTATTTCTAAGACCCTT						
55		Q K K S I Q F H W K N S N Q I K I L G N -						

50

B BS
 BS SC
 AP TR
 N1 NF
 22 11
 5

D M H
 D N I
 E L F
 1 1 1

S
 T
 Y
 1

AGAGCCCCCTGGTAGTAGCCCCCTCAGTGCAATGTAGGAGTCCAAGGGGTAAAAACATAC
 541 ----- 600
 TCTCGGGGGGACCATCATCGGGGAGTCACGTTACATCCTCAGGTTCCCCATTTTGTATG
 10 S P P G S S P S V Q C R S P R G K N I Q -

N BBH S B BS
 M MD ASP A BSSGSC S B N SC
 B ND LPV L APTIAR T A L TR
 O LE UBU U N1NACF X N A NF
 2 11 122 1 221111 1 1 4 11
 15

AGGGGGGGAAGACCCTCTCCGTGTCTCAGCTGGAGCTCCAGGATAGTGGCACCTGGACAT
 20 601 ----- 660
 TCCCCCCTTCTGGGAGAGGCACAGAGTCGACCTCGAGGTCCTATCACCGTGGACCTGTA
 G G K T L S V S Q L E L Q D S G T W T C -

N
 NS
 LP
 AH
 31
 25

M
 B
 O
 2

NV A
 HA L
 EE U
 11 1

GCACTGTCTTGCAGAACCAGAAGAAGGTGGAGTTCAAAATAGACATCGTGGTGCTAGCTT
 30 661 ----- 720
 CGTGACAGAACGTCTTGGTCTTCTCCACCTCAAGTTTTATCTGTAGCACCACGATCGAA
 T V L Q N Q K K V E F K I D I V V L A F -

HS M M
 AT N N
 EU L L
 31 1 1
 35

TCCAGAAGGCCTCCAGCATAGTCTATAAGAAAGAGGGGGAACAGGTGGAGTTCTCCTTCC
 40 721 ----- 780
 AGGTCTTCCGGAGGTTCGTATCAGATATTCTTTCTCCCCCTTGTCCACCTCAAGAGGAAGG
 Q K A S S I V Y K K E G E Q V E F S F P -

A A M
 L L N
 U U L
 1 1 1
 45

CACTCGCCTTTACAGTTGAAAAGCTGACGGGCAGTGGCGAGCTGTGGTGGCAGGCGGAGA
 50 781 ----- 840
 GTGAGCGGAAATGTCAACTTTTCGACTGCCCGTCACCGCTCGACACCACCGTCCGCTCT
 L A F T V E K L T G S G E L W W Q A E R -

5
 H M FM A M
 P N LN U B
 H L ML 3 O
 1 1 11 A 2
 841 GGGCTTCCTCCTCCAAGTCTTGGATCACCTTTGACCTGAAGAACAAGGAAGTGTCTGTAA 900

 CCCGAAGGAGGAGGTT CAGAACCTAGTGGAACTGGACTTCTTGTTCTTCACAGACATT
 10
 A S S S K S W I T F D L K N K E V S V K -
 15
 B BS PS
 SM SCADNPAD A A H
 TA TRVRLUUD L L P
 EE NFAAAM9E U U H
 23 11224161 1 1 1
 20
 AACGGGTTACCCAGGACCCTAAGCTCCAGATGGGCAAGAAGCTCCCGCTCCACCTCACCC 960

 TTGCCCAATGGGTCTCTGGGATTCTGAGGTCTACCCGTTCTTCGAGGGCGAGGTGGAGTGGG
 R V T Q D P K L Q M G K K L P L H L T L -
 25
 BS BSS
 M SC HS D M H SCAHM
 N TR AT D N P TRUAN
 L NF EU E L H NF9EL
 1 11 31 1 1 1 11631
 30
 TGCCCCAGGCCTTGCCCTCAGTATGCTGGCTCTGGAAACCTCACCTTGCCCTTGAAGCGA 1020

 ACGGGGTCCGGAACGGAGTCATACGACCGAGACCTTTGGAGTGGGACCGGGAACCTTCGCT
 P Q A L P Q Y A G S G N L T L A L E A K -
 35
 40
 S BS
 F SC H D A
 A TR P D L
 N NF H E U
 1 11 1 1 1
 45
 AAACAGGAAAGTTGCATCAGGAAGTGAACCTGGTGGTGATGAGAGCCACTCAGCTCCAGA 1080

 TTTGTCCTTTCAACGTAGTCCTTCACTTGGACCACCACTACTCTCGGTGAGTCGAGGTCT
 T G K L H Q E V N L V V M R A T Q L Q K -
 50
 55

M PS S
 N ADNNPA DF AM DE A
 L VRLUU DA LN DS L
 1 AAAAM9 EN UL EP U
 224416 11 11 11 1
 // // //
 1081 AAAATTTGACCTGTGAGGTGTGGGGACCCACCTCCCCTAAGCTGATGCTGAGCTTGAAC 1140

 10 TTTTAAACTGGACACTCCACACCCCTGGGTGGAGGGGATTGACTACGACTCGAACTTTG
 N L T C E V W G P T S P K L M L S L K L -
 M T H M DW
 15 N A P N DS
 L Q A L ET
 1 1 2 1 12
 /
 20 1141 TGGAGAACAAGGAGGCAAAGGTCTCGAAGCGGGAGAAGCCGGTGTGGGTGCTGAACCCCTG 1200

 ACCTCTTGTTCTCCGTTTCCAGAGCTTCGCCCTCTTCGCCACACCCACGACTTGGGAC
 E N K E A K V S K R E K P V W V L N P E -
 25 F D M I A PS H
 O D A N V ADPA I
 K E E F A VRUU N
 1 1 3 1 1 AAM9 F
 2216 1
 ///
 30 1201 AGGCGGGGATGTGGCAGTGTCTGCTGAGTGACTCGGCACAGGTCCTGCTGGAATCCAACA 1260

 TCCGCCCCCTACACCGTCACAGACGACTCACTGAGCCCTGTCCAGGACGACCTTAGGTTGT
 A G M W Q C L L S D S G Q V L L E S N I -
 S SA BHF BS B
 40 ANA HNCP SGNMAANXA SH
 VLU PCRA PIUNMULHV PP
 AA9 AIFL 1ADLH3A0A 1H
 236 2111 21211A421 21
 // // // //
 1261 TCAAGGTTCTGCCCACATGGTCCACCCCGGTGCACGCGGATCCCGAGGGTGAGTGTCGCC 1320

 45 AGTTCCAAGACGGGTGTACCAGGTGGGGCCACGTGCGCCTAGGGCTCCCACTCACACGGG
 K V L P T W S T P V H A D P E
 50
 55

MF BS S S S A M M
 AD SC F DHNA HNC F A B
 EK TR A RALU PCR F A B
 11 NF N AEA9 AIF L E 0
 / 11 1 2346 211 3 2 2

1321 TAGAGTAGCCTGCATCCAGGGACAGGCCCCAGCCGGGTGCTGACACGTCCACCTCCATCT 1380
 ATCTCATCGGACGTAGGTCCCTGTCCGGGGTCGGCCACGACTGTGCAGGTGGAGGTAGA

M D M BS S M S
 N D N SC M ANA M M S
 L E L NF 0 AA9 0 L Y
 1 1 1 11 2 246 2 1 1

1381 CTTCTCAGCACCTGAACTCCTGGGGGGACCGTCAGTCTTCTCTTCCCCCAAACCCA 1440
 GAAGGAGTCGTGGACTTGAGGACCCCCCTGGCAGTCAGAAGGAGAAGGGGGGTTTGGGT

A P E L L G G P S V F L F P P K P K -

S SS N
 AN M HMANNAC DM M NS
 UL N PNVLUR DS A LP
 3A L ALAIA9F ET E AH
 A3 1 2121461 12 3 31 2

1441 AGGACACCCTCATGATCTCCCGGACCCCTGAGGTACATGCGTGGTGGTGGACGTGAGCC 1500
 TCCTGTGGGAGTACTAGAGGGCCTGGGGACTCCAGTGTACGCACCACCACCTGCACTCGG

D T L M I S R T P E V T C V V V D V S H -

M DM M RM M
 N DS B SA N
 L ET 0 AE L
 1 12 2 12 1

1501 ACCAAGACCCTGAGGTCAAGTTCAACTGGTACGTGGACGGCGTGGAGGTGCATAATGCCA 1560
 TGCTTCTGGGACTCCAGTTCAAGTTGACCATGCACCTGCCGCACCTCCACGTATTACGGT

E D P E V K F N W Y V D G V E V H N A K -

F FN S
 M N NSS R M R HNC HH
 N U UPA S A S PCR GP
 L 4 DBC A E A AIF AH
 1 H 222 1 2 1 211 11

1561 AGACAAAGCCGCGGGAGGAGCAGTACAACAGCACGTACCGGGTGGTCAAGCTCCTCACCG 1620
 TCTGTTTCGGCGCCCTCCTCGTCATGTTGTCTGTCATGCCCCACCACTCGCAGGAGTGGC

T K P R E E Q Y N S T Y R V V S V L T V -

5 M N L 1 BS SC TR NF 11 R S A 1

10 TCCTGCACCAAGGACTGGCTGAATGGCAAGGAGTACAAGTGCAAGGTCTCCAACAAAGCCC 1621 1680
 AGGACGTGGTCCTGACCGACTTACCGTTCCTCATGTTACGTTCCAGAGGTGTTTCGGG
 L H Q D W L N G K E Y K C K V S N K A L -

15 P S S
 ADNNPMA A
 VRLLUNU U
 AAAAML9 9
 2244116 6

20 TCCCAGCCCCCATCGAGAAAACCATCTCCAAAGCCAAAGGTGGGACCCGTGGGGTGGCAG 1681 1740
 AGGGTCGGGGGTAGCTCTTTTGGTAGAGGTTTCGGTTTCCACCCTGGGCACCCACGCTC
 P A P I E K T I S K A K

25 H M N HHN S DM M N R
 A N L APA BSAH D N A P S
 E L A EAE GFUA E L E B A
 3 1 3 321 1163 1 1 3 2 1

30 GGCCACATGGACAGAGGCCCGGCTCGGCCCCACCCTCTGCCCTGAGAGTGACCGCTGTACCA 1741 1800
 CCGGTGTACCTGTCTCCGGCCGAGCCGGGTGGGAGACGGGACTCTCACTGGCGACATGGT

35 M N A B R F SS
 N U V B S O AHNNCC
 L 4 A V A K VPCCRR
 1 H 1 1 1 1 121111

40 ACCTCTGTCTACAGGGCAGCCCCGAGAACCACAGGTGTACACCCTGCCCCATCCCGGG 1801 1860
 TGGAGACAGGATGTCCCGTCGGGGCTCTTGGTGTCCACATGTGGGACGGGGGTAGGGCCC
 G Q P R E P Q V Y T L P P S R D -

45

50

55

5
 S A F BS BS B
 M L O SC SC S
 A U K TR TR P
 1 1 1 11 11 1
 / / /
 1861 ATGAGCTGACCAAGAACCAGGTCAGCCTGACCTGCCTGGTCAAAGGCTTCTATCCCAGCG 1920
 10 TACTCGACTGGTTCTTGGTCCAGTCGGACTGGACGGACCAGTTTCCGAAGATAGGGTCGC
 E L T K N Q V S L T C L V K G F Y P S D -
 15 F N H B
 U P B
 4 A V
 H 2 1
 1921 ACATCGCCGTGGAGTGGGAGAGCAATGGGCAGCCGGAGAACAACACTACAAGACCACGCCTC 1980
 20 TGTAGCGGCACCTCACCTCTCGTTACCCGTCGGCCTCTTGTGATGTTCTGGTGCGGAG
 I A V E W E S N G Q P E N N Y K T T P P -
 25 H M N H M A B
 M I M N P N L S
 N N B L P N L P
 L F O A H L U M
 1 1 2 4 1 1 1
 1981 CCGTGCTGGACTCCGACGGCTCCTTCTTCTCTACAGCAAGCTCACCGTGGACAAGAGCA 2040
 30 GGCACGACCTGAGGCTGCCGAGGAAGAAGGAGATGTCGTTGAGTGGCACCTGTTCTCGT
 V L D S D G S F F L Y S K L T V D K S R -
 35 F S
 M M MBX NF M N N
 U B ABM LA N S L
 40 EVN AN L I A
 H2 211 31 1 1 3
 / /
 2041 GGTGGCAGCAGGGGAACGTCTTCTCATGCTCCGTGATGCATGAGGCTCTGCACAACCACT 2100
 CCACCGTCGTCCCCCTTGCAGAAGAGTACGAGGCACTACGTA TCCGAGACGTGTTGGTGA
 W Q Q G N V F S C S V M H E A L H N H Y -
 45 S
 M M HNC CXH
 B N PCR FMA
 O L AIF RAE
 50 2 1 211 133
 / /
 2101 ACACGCAGAAGAGCCTCTCCCTGTCTCCGGGTAAATGAGTGGCAGGCGCG 2150
 TGTGCGTCTTCTCGAGAGGGACAGAGGCCATTTACTCAGCTGCCCGC
 55 T Q K S L S L S P G K .

Example 2: Preparation of the Fusion Proteins from Supernatants of COS Cells

COS cells grown in DME medium supplemented with 10% Calf Serum and gentamicin sulfate at 15 $\mu\text{g/ml}$ were split into DME medium containing 10% NuSerum (Collaborative Research) and gentamicin to give 50% confluence the day before transfection. The next day, CsCl purified plasmid DNA was added to a final concentration of 0.1 to 2.0 $\mu\text{g/ml}$ followed by DEAE Dextran to 400 $\mu\text{g/ml}$ and chloroquine to 100 μM . After 4 hours at 37°C, the medium was aspirated and a 10% solution of dimethyl sulfoxide in phosphate buffered saline was added for 2 minutes, aspirated, and replaced with DME/10% Calf Serum. 8 to 24 hours later, the cells were trypsinized and split 1:2.

For radiolabeling, the medium was aspirated 40 to 48 hours after transfection, the cells washed once with phosphate buffered saline, and DME medium lacking cysteine or methionine was added. 30 minutes later, ^{35}S -labeled cysteine and methionine were added to final concentrations of 30-60 μCi and 100-200 μCi respectively, and the cells allowed to incorporate label for 8 to 24 more hours. The supernatants were recovered and examined by electrophoresis on 7.5% polyacrylamide gels following denaturation and reduction, or on 5% polyacrylamide following denaturation without reduction. The CD4B γ 1 protein gave the same molecular mass with or without reduction, while the CD4E γ 1 and CD4H γ 1 fusion proteins showed molecular masses without reduction of twice the mass observed with reduction, indicating that they formed dimer structures. The CD4 IgM fusion proteins formed large multimers beyond the resolution of the gel system without reduction, and monomers of the expected molecular mass with reduction.

Unlabeled proteins were prepared by allowing the cells to grow for 5 to 10 days post transfection in DME medium containing 5% NuSerum and gentamicin as above. The supernatants were harvested, centrifuged, and purified by batch adsorption to either protein A trisacryl, protein A agarose, goat anti-human IgG antibody agarose, rabbit anti-human IgM antibody agarose, or monoclonal anti-CD4 antibody agarose. Antibody agarose conjugates were prepared by coupling purified antibodies to cyanogen bromide activated agarose according to the manufacturer's recommendations, and using an antibody concentration of 1 mg/ml. Following batch adsorption by shaking overnight on a rotary table, the beads were harvested by pouring into a sintered glass funnel and washed a few times on the funnel with phosphate buffered saline containing 1% Nonidet P40 detergent. The beads were removed from the funnel and poured into a small disposable plastic column (Quik-Sep QS-Q column, Isolab), washed with at least 20 column volumes of phosphate buffered saline containing 1% Nonidet P40, with 5 volumes of 0.15 N NaCl, 1 mM EDTA (pH 8.0), and eluted by the addition of either 0.1 M acetic acid, 0.1 M acetic acid containing 0.1 M NaCl, or 0.25 M glycine-HCl buffer, pH 2.5.

Example 3: Blockage of Syncytium Formation by the Fusion Proteins

Purified or partially purified fusion proteins were added to HPB-ALL cells infected 12 hours previously with a vaccinia virus recombinant encoding HIV envelope protein. After incubation for 6-8 more hours, the cells were washed with phosphate buffered saline, fixed with formaldehyde, and photographed. All of the full-length CD4 immunoglobulin fusion proteins showed inhibition of syncytium formation at a concentration of 20 $\mu\text{g/ml}$ with the exception of the 4H γ 1 protein, which was tested only at 5 $\mu\text{g/ml}$ and showed partial inhibition of syncytium formation under the same conditions.

Example 4: Chromium Release Cytolysis Assay

The purified fusion proteins were examined for ability to fix complement in a chromium release assay using vaccinia virus infected cells as a model system. Namalwa (B cell) or HPB-ALL (T cell) lines were infected with vaccinia virus encoding HIV envelope protein, and 18 hours later were radiolabeled by incubation in 1 mCi/ml sodium ^{51}Cr chromate in phosphate buffered saline for 1 hour at 37°C. The labeled cells were centrifuged to remove the unincorporated chromate, and incubated in microtiter wells with serial dilutions of the CD4 immunoglobulin fusion proteins and rabbit complement at a final concentration of 40%. After 1 hour at 37°C, the cells were mixed well, centrifuged, and the supernatants counted in a gamma-ray counter. No specific release could be convincingly documented.

Example 5: Binding of the CD4E γ 1 Protein to Fc Receptors

Purified CD4E γ 1 fusion protein was tested for its ability to displace radiolabeled human IgG1 from human Fc receptors expressed on COS cells in culture. The IgG1 was radiolabeled with sodium ¹²⁵I using 1 mCi of iodide, 100 μ g of IgG1, and two iodobeads (Pierce). The labeled protein was separated from unincorporated counts by passage over a Sephadex G25 column equilibrated with phosphate buffered saline containing 0.5 mM EDTA and 5% nonfat milk. Serial dilutions of the CD4E γ 1 fusion protein or unlabeled IgG1 were prepared and mixed with a constant amount of radiolabeled IgG1 tracer. After incubation with COS cells bearing the FcRI and FcRII receptors at 4 °C for at least 45 minutes in a volume of 20 μ l, 200 μ l of a 3:2 mixture of dibutyl to dioctyl phthalates were added, and the cells separated from the unbound label by centrifugation in a microcentrifuge for 15 to 30 seconds. The tubes were cut with scissors, and the cell pellets counted in a gamma-ray counter. The affinity of the CD4E γ 1 protein for receptors was measured in parallel with the affinity of the authentic IgG1 protein, and was found to be the same, within experimental error.

Example 6: Stable Expression of the Fusion Construct pCD4E γ 1 in Baby Hamster Kidney Cells

Twenty-four hours before transfection, 0.5×10^6 baby hamster kidney cells (BHK; ATCC CCL10) were seeded in a 25 cm² culture flask in Dulbecco's modified Eagle's medium (DMEM) containing 10% of fetal calf serum (FCS). The cells were cotransfected with a mixture of the plasmids pCD4E γ 1 (20 μ g), pSV2dhfr (5 μ g; Lee et al., *Nature* 294:228-232 (1981)) and pRMH140 (5 μ g, Hudziak et al., *Cell* 31:137-146 (1982)) according to a modified calcium phosphate transfection technique as described in Zettlmeissl et al. (*Behring Inst. Res. Comm.* 82:26-34 (1988)). 72 h post-transfection, cells were split 1:3 to 1:4 (60 mm culture dishes) and resistant colonies were selected in DMEM medium containing 10% FCS, 400 μ g/ml G418 (Geneticin, Gibco) and 1 μ M methotrexate (selection medium). The medium was changed twice a week. The resistant colonies (40-100/transfection) appeared 10-15 day post-transfection and were further propagated either as a mixture of clones (i.e., BHK-NK1) or as individually isolated clones. For the determination of the relative expression levels, clone mixtures or individual clones were grown to confluency in T25 culture flasks, washed twice with protein-free DMEM medium, and incubated for 24 h with 5 ml protein-free DMEM medium. These media were collected and subjected to a human IgG specific ELISA in order to determine the relative expression levels of the CD4-IgG1 fusion protein CD4E γ 1. For further analysis an individual clone (BHK-UC3) was chosen due to its high relative expression levels.

Example 7: Detection of the CD4E γ 1 Protein in Culture Supernatants

For ³⁵S methionine labeling of cells, the clone BHK-UC3 and untransfected BHK cells (control) were grown to confluency in T25 culture flasks and subsequently incubated for two hours in HamF12 medium without methionine. Labeling was achieved by incubating 24 h in 2.5 ml of the same medium containing 100 μ Ci ³⁵S methionine (1070 Ci/mmol, Amersham). For the preparation of cell lysates, the labeled cells were harvested in 1 ml of phosphate buffered saline, pH 7.2 (PBS) and lysed by repetitive freezing and thawing. Cleared lysates (after centrifugation 20000 rpm, 20 min) and culture supernatants were incubated with Protein A-Sepharose (Pharmacia) and the bound material was analyzed on a 10% SDS-Protein A-Sepharose (Pharmacia) and the bound material was analyzed on a 10% SDS-gel according to Laemmli (*Nature* 227:680-685 (1970)), which was subsequently autoradiographed. A specific band of about 80 kDa can be detected only in the supernatant of clone BHK-UC3, which is absent in the lysate of clone BHK-UC3 and in the respective controls.

Example 8: Purification of the Protein CD4E γ 1 from Culture Supernatants

In order to demonstrate that the fusion protein coded by the plasmid pCD4E γ 1 can be obtained in high quantities, the clone BHK-UC3 was grown in 1750 cm² roller bottles in selection medium (500 ml). Confluent monolayers were washed twice with protein-free DMEM medium (200 ml) and further incubated for 48 h with protein-free DMEM medium (500 ml). The conditioned culture supernatants (1-2 l) and respective supernatants from untransfected BHK cells were cleared by centrifugation (9000 rpm, 30 min) and microfiltered through a 0.45 μ m membrane (Nalgene). After addition of 1% (v/v) of 1.9 M Tris-HCl buffer,

pH 8.6, the conditioned medium was absorbed to a Protein A-Sepharose column equilibrated with 50 mM Tris-HCl pH 8.6 buffer containing 150 mM NaCl (4°C). The loaded column was washed with 10 column volumes of equilibration buffer. Elution of the CD4-IgG1 fusion protein CD4E γ 1 was achieved with 0.1 M sodium citrate buffer, pH 3, followed by immediate neutralization of the column efflux to pH 8 by Tris-base.

The peak fractions were pooled, and the pool was analyzed on a Coomassie blue stained SDS-gel resulting in a band of the expected size (80 KDa), and which reacted with a polyclonal anti-human IgG heavy chain antibody and a mouse monoclonal anti-CD4 antibody (BMA040, Behringwerke) in Western Blots. The yields of purified fusion proteins obtained by the given procedure is 5-18 mg/24 h/l culture supernatant. The respective value for a BHK clone mixture (about 80 resistant clones; BHK-NK1) as described above was 2-3 mg/24 h/l.

Example 9: Physical and Biological Characterization of the CD4E γ 1 Fusion Protein

As proven by SDS-electrophoresis on 10-15% gradient gels (Phast-System, Pharmacia) under non-reductive conditions, the CD4E γ 1 fusion protein migrates at the position of a homodimer (about 160 KDa) like a non-reduced mouse monoclonal antibody. This result is supported by analytical equilibrium ultracentrifugation, where the fusion protein behaves as a homogeneous dimeric molecule of about 150 KDa. The absorbance coefficient of the protein was determined as $A_{280} = 18 \text{ cm}^2/\text{mg}$ using the quantitative protein determination according to Bradford (Anal. Biochem. 72:248-254 (1976)).

The CD4E γ 1-fusion protein shows specific complex formation with a solubilized β gal-gp120 fusion protein (pMB1790; Broker et al., Behring Inst. Res. Commun. 82:338-348 (1988)) expressed in *E. coli*. In this protein (110 KDa), a major part of the HIV gp120 protein (Val₄₉-Trp₆₄₆) is fused to β -galactosidase (amino acids 1-375). In a control experiment a 67-KDa β -gal-HIV 3'orf fusion protein (β gal1-375; 3'orf Pro14-Asp123) showed no complex formation. In these experiments, the CD4E γ 1-protein was incubated with the respective fusion protein in molar ratios of about 5:1. The complex was isolated by binding to Protein A-Sepharose and the Protein A-Sepharose bound proteins--together with relevant controls--were analyzed on 10-15% gradient SDS-gels (Phast-System, Pharmacia).

The CD4E γ 1 fusion protein binds to the surface of HIV (HIV1/HTLV-IIIb) infected cultured T4-lymphocytes as determined by direct immunofluorescence with fluorescein-isothiocyanate (FITC) labeled CD4E γ 1 protein. It blocks syncytia formation in cultured T4-lymphocytes upon HIV infection (0.25 TCID₅₀/cell) at a concentration of 10 $\mu\text{g}/\text{ml}$. Furthermore, HIV-infected cultured T4-lymphocytes (subclone of cell line H9) are selectively killed upon incubation with CD4E γ 1 in the presence or absence of complement: To a highly (>50%) HIV infected culture of T4-lymphocytes (10^6 cells/ml) 50, 10 or 1 $\mu\text{g}/\text{ml}$ CD4E γ 1 fusion protein was added in the presence or absence of guinea pig complement. Cells were observed for specific killing by the fusion protein, which is defined by the percentage of killed cells after 3 days in relation to viable cells in the culture at the beginning of the experiment corrected by the values for unspecific killing observed in control cultures, lacking the CD4E γ 1 fusion protein (Table 5, Experiment I). Surprisingly, addition of CD4E γ 1 protein to the infected T4 cells in the absence of complement resulted in similar specific killing rates as in the presence of complement (Table 5, Experiment II). This result demonstrates a complement independent cytolytic effect of CD4E γ 1 on HIV infected T-lymphocytes in culture.

Table 5

No. Experiment	Assay System	Specific Killing (%)
I	non-infected T4-cells + 50 $\mu\text{g}/\text{ml}$ CD4E γ 1 + Compl.	0.7
	infected T4-cells + 50 $\mu\text{g}/\text{ml}$ CD4E γ 1 + Compl.	35.1
	infected T4-cells + 10 $\mu\text{g}/\text{ml}$ CD4E γ 1 + Compl.	25.1
	infected T4-cells + 1 $\mu\text{g}/\text{ml}$ CD4E γ 1 + Compl.	25
II	infected T4-cells + 10 $\mu\text{g}/\text{ml}$ CD4E γ 1 + Compl.	49.9
	infected T4-cells + 10 $\mu\text{g}/\text{ml}$ CD4E γ 1 + Compl.	69.4

Having now fully described this invention, it will be appreciated by those skilled in the art that the same can be performed with any wide range of equivalent parameters of composition, conditions, and methods of preparing such fusion proteins without departing from the spirit or scope of the invention or any embodiment thereof.

Claims

1. A fusion protein gene comprising 1) the DNA sequence of CD4, or fragment thereof which binds to HIV gp120, and 2) the DNA sequence of an immunoglobulin heavy chain, characterized in that the DNA sequence which encodes the variable region of said immunoglobulin chain has been replaced with the DNA sequence which encodes CD4, or said gp120 binding fragment thereof.

2. The fusion protein gene of claim 1, wherein the DNA sequence which encodes said fragment of CD4 comprises the following DNA sequence:

10

CAATGAACCGGG

-+----- 120

GTTACTTGGCCC

15

GAGTCCCTTTTAGGCACTTGCTTCTGGTGCTGCAACTGGCGCTCCTCCCAGCAGCCACTC

121 -----+-----+-----+-----+-----+----- 180

CTCAGGGGAAAATCCGTGAACGAAGACCACGACGTTGACCGCGAGGAGGCTCGTCGGTGAG

20

AGGGAAAGAAAGTGCTGCTGGGCAAAAAGGGGATACAGTGGAACTGACCTGTACAGCTT

181 -----+-----+-----+-----+-----+----- 240

TCCCTTTCTTTTACCACGACCCGTTTTTTCCCCTATGTCACCTTGACTGGACATGTCGAA

25

CCCAGAAGAAGAGCATACAATTCCACTGGAAAACTCCAACCAGATAAAGATTCTGGGAA

241 -----+-----+-----+-----+-----+----- 300

GGGTCTTCTTCTCGTATGTTAAGGTGACCTTTTGGAGGTGGTCTATTTCTAAGACCTT

30

ATCAGGGCTCCTTCTTAATAAGGTCCATCCAAGCTGAATGATCGCGCTGACTCAAGAA

301 -----+-----+-----+-----+-----+----- 360

TAGTCCCGAGGAAGAATTGATTTCAGGTAGGTTGACTTACTAGCGCGACTGAGTTCTT

35

GAAGCCTTTGGGACCAAGGAACTTCCCCCTGATCATCAAGAATCTTAAGATAGAAGACT

361 -----+-----+-----+-----+-----+----- 420

CTTCGGAAACCCTGGTTCCTTTGAAGGGGGACTAGTAGTTCTTAGAATTCTATCTTCTGA

40

CAGATACTTACATCTGTGAAGTGAGGACCAGAAGGAGGAGGTGCAATTGCTAGTGTTCTG

421 -----+-----+-----+-----+-----+----- 480

GTCTATGAATGTAGACACTTCACCTCCTGGTCTTCCTCCTCCACGTTAACGATCACAAGC

45

GATTGACTGCCAACTCTGACACCCACCTGCTTC

481 -----+-----+-----+-----

50

CTAACTGACGGTTGAGACTGTGGGTGGACGAAG

or a degenerate variant thereof, or the following DNA sequence:

55

CAATGAACCGGG

-+----- 120

GTTACTTGGCCC

5

10

15

20

25

30

35

40

45

50

GAGTCCCTTTTAGGCACTTGCTTCTGGTGCTGCAACTGGCGCTCCTCCCAGCAGCCACTC
 121 -----+-----+-----+-----+-----+----- 180
 CTCAGGGAAAATCCGTGAACGAAGACCACGACGTTGACCGCGAGGAGGGTCGTCGGTGAG
 AGGGAAAGAAAGTGGTGCTGGGCAAAAAGGGGATACAGTGGAAGTACCTGTACAGCTT
 181 -----+-----+-----+-----+-----+----- 240
 TCCCTTTCTTTCACCACGACCCGTTTTTCCCTATGTCACCTTGACTGGACATGTCGAA
 CCCAGAAGAAGAGCATACAATTCCACTGGAAAACTCCAACCAGATAAAGATTCTGGGAA
 241 -----+-----+-----+-----+-----+----- 300
 GGGTCTTCTTCTCGTATGTTAAGGTGACCTTTTTGAGGTTGGTCTATTTCTAAGACCCTT
 ATCAGGGCTCCTTCTTAAGTAAAGGTCCATCCAAGCTGAATGATCGCGCTGACTCAAGAA
 301 -----+-----+-----+-----+-----+----- 360
 TAGTCCCGAGGAAGAATTGATTTCCAGGTAGGTTGCACTTACTAGCGCGACTGAGTTCTT
 GAAGCCTTTGGGACCAAGGAACTTCCCCCTGATCATCAAGAATCTTAAGATAGAAGACT
 361 -----+-----+-----+-----+-----+----- 420
 CTTCCGAAACCCTGGTTCCTTTGAAGGGGGACTAGTAGTTCTTAGAATTCTATCTTCTGA
 CAGATACTTACATCTGTGAAGTGGAGGACCAGAAGGAGGAGGTGCAATTGCTAGTGTTCCG
 421 -----+-----+-----+-----+-----+----- 480
 GTCTATGAATGTAGACACTTCACCTCCTGGTCTTCCTCCTCCACGTTAACGATCACAAGC
 GATTGACTGCCAACTCTGACACCCACCTGCTTCAGGGGCAGAGCCTGACCCTGACCTTGG
 481 -----+-----+-----+-----+-----+----- 540
 CTAAGTACGGTTGAGACTGTGGGTGGACGAAGTCCCCGTCTCGGACTGGGACTGGAACC
 AGAGCCCCCCTGGTAGTAGCCCCCTCAGTGCAATGTAGGAGTCCAAGGGGTAAAAACATAC
 541 -----+-----+-----+-----+-----+----- 600
 TCTCGGGGGGACCATCATCGGGGAGTCACGTTACATCCTCAGGTTCCCCATTTTGTATG
 AGGGGGGAAGACCCTCTCCGTCTCTCAG
 601 -----+-----+-----+-----+-----
 TCCCCCCTTCTGGGAGAGGCACAGAGTC

or a degenerate variant thereof.

55

3. The fusion protein gene of claim 1 or 2, characterized in that said immunoglobulin chain is of the class IgM, IgG1 or IgG3.

4. A fusion protein gene comprising 1) the DNA sequence of CD4, or fragment thereof which binds to HIV gp120, and 2) the DNA sequence of an immunoglobulin light chain, characterized in that the DNA sequence which encodes the variable region of said immunoglobulin light chain has been replaced with the DNA sequence which encodes CD4, or HIV gp120-binding fragment thereof.

5. A fusion protein gene of claim 4, characterized in that the DNA sequence which encodes said fragment of CD4 comprises the following DNA sequence:

```

                                CAATGAACCGGG
                                -+----- 120
                                GTTACTTGGCCC

10
                                GAGTCCCTTTTAGGCACTTGCTTCTGGTGCTGCAACTGGCGCTCCTCCCAGCAGCCACTC
121 -----+-----+-----+-----+-----+----- 180
                                CTCAGGGAAAATCCGTGAACGAAGACCACGACGTTGACCGCGAGGAGGGTCGTCCGGTGAG

15
                                AGGGAAAGAAAGTGGTGCTGGGCCAAAAAGGGGATACAGTGGAAGTACCTGTACAGCTT
181 -----+-----+-----+-----+-----+----- 240
                                TCCCTTTCTTTTACCACGACCCGTTTTTTCCCTATGTCACCTTGACTGGACATGTCCGAA

20
                                CCCAGAAGAAGAGCATACAATTCCACTGGAAAACTCCAACCAGATAAAGATTCTGGGAA
241 -----+-----+-----+-----+-----+----- 300
                                GGGTCTTCTTCTCGTATGTTAAGGTGACCTTTTTGAGGTTGGTCTATTTCTAAGACCCCTT

25
                                ATCAGGGCTCCTTCTTAACTAAAGGTCCATCCAAGCTGAATGATCGCGCTGACTCAAGAA
301 -----+-----+-----+-----+-----+----- 360
                                TAGTCCCGAGGAAGAATTGATTCCAGGTAGGTTGACTTACTAGCGCGACTGAGTTCTT

30
                                GAAGCCTTTGGGACCAAGGAAACTTCCCCCTGATCATCAAGAATCTTAAGATAGAAGACT
361 -----+-----+-----+-----+-----+----- 420
                                CTTTCGGAAACCCTGGTTCCTTTGAAGGGGACTAGTAGTTCTTAGAATTCTATCTTCTGA

35
                                CAGATACTTACATCTGTGAAGTGGAGGACCAGAAGGAGGAGGTGCAATTGCTAGTGTTCG
421 -----+-----+-----+-----+-----+----- 480
                                GTCTATGAATGTAGACACTTCACCTCCTGGTCTTCTCCTCCACGTTAACGATCACAAGC

40
                                GATTGACTGCCAACTCTGACACCCACCTGCTTC
481 -----+-----+-----+-----
                                CTAAGTACGCTTGAGACTGTGGGTGGACGAAG

45

```

50 or a degenerate variant thereof, or the following DNA sequence:

55

CAATGAACCGGG

-+----- 120

GTTACTTGGCCC

5

10

15

20

25

30

35

40

45

50

GAGTCCCTTTTAGGCACTTGCTTCTGGTGCTGCAACTGGCGCTCCTCCCAGCAGCCACTC
121 -----+-----+-----+-----+-----+----- 180
CTCAGGGAAAATCCGTGAACGAAGACCACGACGTTGACCGCGAGGAGGGTCGTCGGTGAG
AGGGAAAGAAAGTGCTGCTGGGCAAAAAGGGGATACAGTGGAAGTACCTGTACAGCTT
181 -----+-----+-----+-----+-----+----- 240
TCCCTTTCTTTTACCACGACCCGTTTTTTCCCCTATGTCACCTTGACTGGACATGTCGAA
CCCAGAAGAAGAGCATACAATTCCACTGGAAAACTCCAACCAGATAAAGATTCTGGGAA
241 -----+-----+-----+-----+-----+----- 300
GGGTCTTCTTCTCGTATGTTAAGGTGACCTTTTTGAGGTTGGTCTATTTCTAAGACCCCTT
ATCAGGGCTCCTTCTTAACTAAAGGTCCATCCAAGCTGAATGATCGCGCTGACTCAAGAA
301 -----+-----+-----+-----+-----+----- 360
TAGTCCCAGGAAGAATTGATTTCCAGGTAGGTTGACTTACTAGCGGACTGAGTTCTT
GAAGCCTTTGGGACCAAGGAACTTCCCCCTGATCATCAAGAATCTTAAGATAGAAGACT
361 -----+-----+-----+-----+-----+----- 420
CTTCGGAACCCCTGGTTCCTTTGAAGGGGGACTAGTAGTTCCTTAGAATTCTATCTTCTGA
CAGATACTTACATCTGTGAAGTGGAGGACCAGAAGGAGGAGGTGCAATTGCTAGTGTTCC
421 -----+-----+-----+-----+-----+----- 480
GTCTATGAATGTAGACACTTCACCTCCTGGTCTTCCTCCTCCACGTTAACGATCACAAGC
GATTGACTGCCAACTCTGACACCCACCTGCTTCAGGGGCAGAGCCTGACCCTGACCTTGG
481 -----+-----+-----+-----+-----+----- 540
CTAACTGACGGTTGAGACTGTGGGTGGACGAAGTCCCCGTCTCGGACTGGGACTGGAACC
AGAGCCCCCCTGGTAGTACCCCTCAGTGCAATGTAGGAGTCCAAGGGGTAAAAACATAC
541 -----+-----+-----+-----+-----+----- 600
TCTCGGGGGGACCATCATCGGGGAGTCACGTTACATCCTCAGGTTCCCCATTTTTGTATG
AGGGGGGGAAGACCCTCTCCGTGTCTCAG
601 -----+-----+-----+-----+-----+-----
TCCCCCCCCTTCTGGGAGAGGCACAGAGTC

or a degenerate variant thereof.

55

6. A vector comprising the fusion protein gene of claim 1, preferably having the identifying characteristics of pCD4H γ 1, which has been deposited under Accession No. 67611, or pCD4M μ , which has been deposited under Accession No. 67608, or of pCD4P μ , which has been deposited under Accession No. 67609, or of pCD4E γ 1, which has been deposited under Accession No. 67610, all in E. coli at the ATCC under the terms of the Budapest Treaty.
7. A vector comprising the fusion protein gene of claim 4.
8. A host transformed with the vector of claim 6 or 7.
9. The host of claim 8 which expresses an immunoglobulin light chain together with the expression product of said fusion protein gene to give an immunoglobulin-like molecule which binds to gp120 or an immunoglobulin heavy chain together with the expression product of said fusion protein gene to give an immunoglobulin-like molecule which binds to HIV or SIV gp120.
10. The host of claim 9, wherein said immunoglobulin heavy chain is of the immunoglobulin class IgM, IgG1 or IgG3.
11. A method of producing a fusion protein comprising CD4, or fragment thereof which binds to gp120, and immunoglobulin heavy chain, wherein the variable region of the immunoglobulin chain has been substituted with CD4, or fragment thereof which binds to HIV or SIV gp120, characterized by cultivating in a nutrient medium under protein-producing conditions, a host strain transformed with the vector of claim 6, said vector further comprising expression signals which are recognized by said host strain and direct expression of said fusion protein, and recovering the fusion protein so produced.
12. The method of claim 11, wherein said host strain is a myeloma cell line which produces immunoglobulin light chains and said fusion protein comprises an immunoglobulin heavy chain of the class IgM, IgG1 or IgG3, wherein an immunoglobulin-like molecule comprising said fusion protein is produced.
13. A method of producing a fusion protein comprising CD4, or fragment thereof which binds to gp120, and an immunoglobulin light chain, wherein the variable region of the immunoglobulin chain has been substituted with CD4, or fragment thereof which binds to HIV or SIV gp120, characterized by cultivating in a nutrient medium under protein-producing conditions, a host strain transformed with the vector of claim 7, said vector further comprising expression signals which are recognized by said host strain and direct expression of said fusion protein, and recovering the fusion protein so produced.
14. The method of claim 13, wherein said host produces immuno-globulin heavy chains of the class IgM, IgG1 and IgG3 together with said fusion protein to give an immunoglobulin-like molecule which binds to HIV-gp120.
15. A fusion protein, which is preferably detectably labeled, comprising CD4, or fragment thereof which is capable of binding to HIV or SIV gp120, fused at the C-terminus to a second protein which comprises an immunoglobulin heavy chain of the class IgM, IgG1 or IgG3, wherein the variable region of said heavy chain immunoglobulin has been replaced with CD4, or HIV gp120-binding fragment thereof, and preferably further comprising a therapeutic agent, radiolabel or NMR imaging agent linked to said fusion protein.
16. The fusion proteins CD4H γ 1, CD4M μ , CD4P μ , CD4E γ 1 or CD4B γ 1.
17. An immunoglobulin-like molecule, comprising the fusion protein of claim 15 and an immunoglobulin light chain, preferably further comprising a detectable label, and especially further comprising a therapeutic agent, radiolabel or NMR imaging agent linked to said immunoglobulin-like molecule.
18. A fusion protein comprising CD4, or fragment thereof which binds to HIV gp120, fused at the C-terminus to a second protein comprising an immunoglobulin light chain where the variable region has been deleted, and which fusion protein preferably is detectably labeled, especially further comprising a therapeutic agent, radiolabel or NMR imaging agent linked to said fusion protein.
19. The fusion protein of claim 15, wherein said CD4 fragment comprises the following amino acid sequence:

M N R G

V P F R H L L L V L Q L A L L P A A T Q
 G K K V V L G K K G D T V E L T C T A S
 Q K K S I Q F H W K N S N Q I K I L G N
 Q G S F L T K G P S K L N D R A D S R R
 S L W D Q G N F P L I I K N L K I E D S
 D T Y I C E V E D Q K E E V Q L L V F G
 L T A N S D T H L L Q

or the following amino acid sequence:

M N R G

V P F R H L L L V L Q L A L L P A A T Q
 G K K V V L G K K G D T V E L T C T A S
 Q K K S I Q F H W K N S N Q I K I L G N
 Q G S F L T K G P S K L N D R A D S R R
 S L W D Q G N F P L I I K N L K I E D S
 D T Y I C E V E D Q K E E V Q L L V F G
 L T A N S D T H L L Q G Q S L T L T L E
 S P P G S S P S V Q C R S P R G K N I Q
 G G K T L S V S Q

20. An immunoglobulin-like molecule comprising the fusion protein of claim 18 and an immunoglobulin heavy chain of the class IgM, IgG1 or IgG3, preferably further comprising a detectable label, and especially further comprising a therapeutic agent, radiolabel or NMR imaging agent linked to said immunoglobulin-like molecule.

21. A complex comprising the fusion protein of claim 15 or 18 and HIV or SIV gp120.

22. The complex of claim 21, wherein said gp120 is a part of an HIV or SIV, is expressed on the surface of an HIV or SIV-infected cell or is present in solution.

23. A method for the detection of HIV or SIV gp120 in a sample, characterized by

(a) contacting a sample suspected of containing HIV or SIV gp120 with the fusion protein of claim 15 or 18, and

(b) detecting whether a complex is formed, said fusion protein preferably being detectably labeled.

Claims for the following Contracting State: GR

1. A vector comprising a fusion protein gene comprising 1) the DNA sequence of CD4, or fragment thereof which binds to HIV gp120, and 2) the DNA sequence of an immunoglobulin heavy chain, characterized in that the DNA sequence which encodes the variable region of said immunoglobulin chain has been replaced with the DNA sequence which encodes CD4, or said gp120 binding fragment thereof.

2. The vector of claim 1, having the identifying characteristics of pCD4H γ 1, which has been deposited in *E. coli* at the ATCC under the terms of the Budapest Treaty under Accession No. 67611.

3. The vector of claim 1, having the identifying characteristics of pCD4M μ , which has been deposited in *E. coli* at the ATCC under the terms of this Budapest Treaty under Accession No. 67608.

4. The vector of claim 1, having the identifying characteristics of PCD4P μ , which has been deposited in *E. coli* at the ATCC under the Budapest Treaty under Accession No. 67609.

5. The vector of claim 1, having the identifying characteristics of PC4E γ 1, which has been deposited in E. coli at the ATCC under the terms of the Budapest Treaty under Accession No. 67610.

6. A vector comprising a fusion protein gene characterized by 1) the DNA sequence of CD4, or fragment thereof which binds to HIV gp120, and 2) the DNA sequence of an immunoglobulin light chain, wherein the DNA sequence which encodes the variable region of said immunoglobulin light chain has been replaced with the DNA sequence which encodes CD4, or HIV gp120-binding fragment thereof.

7. A host transformed with the vector of claim 1.

8. The host of claim 7 which expresses an immunoglobulin light chain together with the expression product of said fusion protein gene to give an immunoglobulin-like molecule which binds to gp120.

9. A host transformed with the vector of claim 6.

10. The host of claim 6 which expresses an immunoglobulin heavy chain together with the expression product of said fusion protein gene to give an immunoglobulin-like molecule which binds to HIV or SIV gp120.

11. The host of claim 10, characterized in that said immunoglobulin heavy chain is of the immunoglobulin class IgM, IgG1 or IgG3.

12. A method of producing a fusion protein comprising CD4, or fragment thereof which binds to gp120, and an immunoglobulin heavy chain, wherein the variable region of the immunoglobulin chain has been substituted with CD4, or fragment thereof which binds to HIV or SIV gp120, characterized by cultivating in a nutrient medium under protein-producing conditions, a host strain transformed with the vector of claim 1, said vector further comprising expression signals which are recognized by said host strain and direct expression of said fusion protein, and recovering the fusion protein so produced.

13. The method of claim 12, characterized in that said host strain is a myeloma cell line which produces immunoglobulin light chains and said fusion protein comprises an immunoglobulin heavy chain of the class IgM, IgG1 or IgG3, wherein an immunoglobulin-like molecule comprising said fusion protein is produced.

14. A method of producing a fusion protein comprising CD4, or fragment thereof which binds to gp120, and an immunoglobulin light chain, wherein the variable region of the immunoglobulin chain has been substituted with CD4, or fragment thereof which binds to HIV or SIV gp120, characterized by cultivating in a nutrient medium under protein-producing conditions, a host strain transformed with the vector of claim 6, said vector further comprising expression signals which are recognized by said host strain and direct expression of said fusion protein, and recovering the fusion protein so produced.

15. The method of claim 14, characterized in that said host produces immunoglobulin heavy chains of the class IgM, IgG1 and IgG3 together with said fusion protein to give an immunoglobulin-like molecule which binds to HIV-gp120.

16. A method for the detection of HIV or SIV gp120 in a sample, characterized by

(a) contacting a sample suspected of containing HIV or SIV gp120 with a fusion protein comprising CD4, or fragment thereof which binds to HIV gp120, and 2) an immunoglobulin heavy chain, wherein the variable region of said immunoglobulin chain has been replaced with CD4, or said gp120 binding fragment thereof, and

(b) detecting whether a complex is formed.

17. The method of claim 16, characterized in that said fusion protein is detectably labeled.

18. A method for the detection of HIV or SIV gp120 in a sample, characterized by

(a) contacting a sample suspected of containing HIV or SIV gp120 with a fusion protein comprising comprising 1) CD4, or fragment thereof which binds to HIV gp120, and 2) an immunoglobulin light chain, wherein the variable region of said immunoglobulin light chain has been replaced with CD4, or HIV gp120-binding fragment thereof, and

(b) detecting whether a complex has formed.

19. The method of claim 18, characterized in that said fusion protein is detectably labeled.

Claims for the following Contracting State: ES

1. A method of producing a fusion protein comprising CD4, or fragment thereof which binds to gp120, and an immunoglobulin heavy chain, wherein the variable region of the immunoglobulin chain has been substituted with CD4, or fragment thereof which binds to HIV or SIV gp120, characterized by cultivating in a nutrient medium under protein-producing conditions, a host strain transformed with a vector comprising a fusion protein gene comprising 1) the DNA sequence of CD4, or fragment thereof which binds to HIV

gp120, and 2) the DNA sequence of an immunoglobulin heavy chain, wherein the DNA sequence which encodes the variable region of said immunoglobulin chain has been replaced with the DNA sequence which encodes CD4, or said gp120 binding fragment thereof, said vector further comprising expression signals which are recognized by said host strain and direct expression of said fusion protein, and recovering the fusion protein so produced.

2. The method of claim 1, characterized in that said vector has the identifying characteristics of pCD4H γ 1, which has been deposited in E. coli at the ATCC under the terms of the Budapest Treaty under Accession No. 67611.

3. The method of claim 1, characterized in that said vector has the identifying characteristics of PCD4M μ , which has been deposited in E. coli at the ATCC under the terms of this Budapest Treaty under Accession No. 67608.

4. The method of claim 1, characterized in that said vector has the identifying characteristics of PCD4P μ , which has been deposited in E. coli at the ATCC under the Budapest Treaty under Accession No. 67609.

5. The method of claim 1, characterized in that said vector has the identifying characteristics of pCD4E γ 1, which has been deposited in E. coli at the ATCC under the terms of the Budapest Treaty under Accession No. 67610.

6. The method of claim 1, characterized in that said host strain is a myeloma cell line which produces immunoglobulin light chains and said fusion protein comprises an immunoglobulin heavy chain of the class IgM, IgG1 or IgG3, wherein an immunoglobulin-like molecule comprising said fusion protein is produced.

7. A method of producing a fusion protein comprising CD4, or fragment thereof which binds to gp120, and an immunoglobulin light chain, wherein the variable region of the immunoglobulin chain has been substituted with CD4, or fragment thereof which binds to HIV or SIV gp120, characterized by cultivating in a nutrient medium under protein-producing conditions, a host strain transformed with a vector comprising a fusion protein gene comprising 1) the DNA sequence of CD4, or fragment thereof which binds to HIV gp120, and 2) the DNA sequence of an immunoglobulin light chain, wherein the DNA sequence which encodes the variable region of said immunoglobulin light chain has been replaced with the DNA sequence which encodes CD4, or HIV gp120-binding fragment thereof, said vector further comprising expression signals which are recognized by said host strain and direct expression of said fusion protein, and recovering the fusion protein so produced.

8. The method any one of claims 1 or 7, characterized in that the DNA sequence which encodes said fragment of CD4 comprises the following DNA sequence:

CAATGAACCGGG

- +----- 120

GTTACTTGGCCC

5

121 GAGTCCCTTTT TAGGCACTTGCTTCTGGTGCTGCAACTGGCGCTCCTCCCAGCAGCCACTC 180
-----+-----+-----+-----+-----+-----
10 CTCAGGGAAAATCCGTGAACGAAGACCACGACGTTGACCGCGAGGAGGGTCGTCGGTGAG
181 AGGGAAAGAAAGTGCTGCTGGGCAAAAAAGGGGATACAGTGGAAGTACCTGTACAGCTT 240
-----+-----+-----+-----+-----+-----
15 TCCCTTTCTTTCACCAACGACCCGTTTTTTCCCCTATGTCACCTTGACTGGACATGTCGAA
241 CCCAGAAGAAGAGCATACAATTCCACTGGAAAACTCCAACCAGATAAAGATTCTGGGAA 300
-----+-----+-----+-----+-----+-----
20 GGGTCTTCTTCTCGTATGTTAAGGTGACCTTTTTGAGGTTGGTCTATTTCTAAGACCTT
301 ATCAGGGCTCCTTCTTAAGTAAAGGTCCATCCAAGCTGAATGATCGCGCTGACTCAAGAA 360
-----+-----+-----+-----+-----+-----
25 TAGTCCCGAGGAAGAATTGATTTCCAGGTAGGTTGACTTACTAGCGCGACTGAGTTCTT
361 GAAGCCTTTGGGACCAAGGAACTTCCCCCTGATCATCAAGAATCTTAAGATAGAAGACT 420
-----+-----+-----+-----+-----+-----
30 CTTCGGAAACCCTGCTTCCTTTGAAGGGGACTAGTAGTTCTTAGAATTCTATCTTCTGA
421 CAGATACTTACATCTGTGAAGTGGAGGACCAGAAGGAGGAGGTGCAATTGCTAGTGTTCG 480
-----+-----+-----+-----+-----+-----
35 GTCTATGAATGTAGACACTTCACCTCCTGGTCTTCCTCCTCCACGTTAACGATCACAAGC
481 GATTGACTGCCAACTCTGACACCCACCTGCTTC
-----+-----+-----+-----
40 CTAAGTACGCGTTGAGACTGTGGGTGGACGAAG

or a degenerate variant thereof.

9. The method of any one of claims 1 or 7, characterized in that said DNA sequence which encodes
said fragment of CD4 comprises the following DNA sequence:

45

50

55

5 CAATGAACCGGG
 -+----- 120
 GTTACTTGGCCC

10 GAGTCCCTTTTAGGCACTTGCTTCTGGTGCTGCAACTGGCGCTCCTCCCAGCAGCCACTC
 121 -----+-----+-----+-----+-----+----- 180
 CTCAGGGAAAATCCGTGAACGAAGACCACGACGTTGACCGCGAGGAGGGTCGTCCGTGAG

15 AGGGAAAGAAAGTGGTGCTGGGCAAAAAAGGGGATACAGTGGAAGTACCTGTACAGCTT
 181 -----+-----+-----+-----+-----+----- 240
 TCCCTTTCTTTTACCACGACCCGTTTTTCCCTATGTCACCTTGACTGGACATGTGCGAA

20 CCCAGAAGAAGAGCATACAATTCCACTGGAAAACTCCAACCAGATAAAGATTCTGGGAA
 241 -----+-----+-----+-----+-----+----- 300
 GGGTCTTCTTCTCGTATGTTAAGGTGACCTTTTTGAGGTTGGTCTATTTCTAAGACCCCTT

25 ATCAGGGCTCCTTCTTAATAAGGTCCATCCAAGCTGAATGATCGCGCTGACTCAAGAA
 301 -----+-----+-----+-----+-----+----- 360
 TAGTCCCAGGAAGAATTGATTTCCAGGTAGGTTGACTTACTAGCGCGACTGAGTTCTT

30 GAAGCCTTTGGGACCAAGGAACTTCCCCCTGATCATCAAGAATCTTAAGATAGAAGACT
 361 -----+-----+-----+-----+-----+----- 420
 CTTCGGAAACCCTGGTTCCTTTGAAGGGGGACTAGTAGTTCTTAGAATTCTATCTTCTGA

35 CAGATACTTACATCTGTGAAGTGGAGGACCAGAAGGAGGAGGTGCAATTGCTAGTGTTCC
 421 -----+-----+-----+-----+-----+----- 480
 GTCTATGAATGTAGACACTTCACCTCCTGGTCTTCCTCCTCCACGTTAACGATCACAAGC

40 GATTGACTGCCAACTCTGACACCCACCTGCTTCAGGGGCAGAGCCTGACCCTGACCTTGG
 481 -----+-----+-----+-----+-----+----- 540
 CTAAGTACGGTTGAGACTGTGGGTGGACGAAGTCCCGTCTCGGACTGGGACTGGAACC

45 AGAGCCCCCCTGGTAGTAGCCCTCAGTGCAATGTAGGAGTCCAAGGGGTAAAAACATAC
 541 -----+-----+-----+-----+-----+----- 600
 TCTCGGGGGGACCATCATCGGGGAGTCACGTTACATCCTCAGGTTCCCCATTTTGTATG

50 AGGGGGGGAAGACCCTCTCCGTGTCTCAG
 601 -----+-----+-----+-----+-----+-----
 TCCCCCCTTCTGGGAGAGGCACAGAGTC

or a degenerate variant thereof.

10. The method of claim 7, characterized in that said host produces immuno-globulin heavy chains of the class IgM, IgG1 and IgG3 together with said fusion protein to give an immunoglobulin-like molecule which binds to HIV-gp120.



Europäisches Patentamt
European Patent Office
Office européen des brevets

Publication number:

0 325 262
A3

EUROPEAN PATENT APPLICATION

Application number: 89100913.6

Date of filing: 20.01.89

Int. Cl.⁵: **C12N 15/00, C12P 21/02,**
A61K 39/395, G01N 33/569,
//A61K39/21

Priority: 22.01.88 US 147351

Date of publication of application:
26.07.89 Bulletin 89/30

Designated Contracting States:
AT BE CH DE ES FR GB GR IT LI LU NL SE

Date of deferred publication of the search report:
16.05.90 Bulletin 90/20

Applicant: **THE GENERAL HOSPITAL**
CORPORATION
55 Fruit Street
Boston MA 02114(US)

Inventor: **Brian, Seed, Dr.**
47A Joy Street
Boston MA 02114(US)

Representative: **Klein, Otto, Dr. et al**
Hoechst AG Zentrale Patentabteilung
Postfach 80 03 20
D-6230 Frankfurt am Main 80(DE)

Cloned genes encoding IG-CD4 fusion proteins and the use thereof.

Fusion proteins of immunoglobulins of the IgM, IgG1 or IgG3 class, wherein the variable region of the light or heavy chain has been replaced with CD4 or fragments thereof capable of binding to gp120 or immunoglobulin-like molecules comprising such fusion proteins together with an immunoglobulin light or heavy chain can be administered to an animal suffering from HIV or SIV infection. They also are useful in assays for HIV or SIV comprising contacting a sample suspected of containing HIV or SIV gp120 with the immunoglobulin-like molecule or fusion protein, and detecting whether a complex is formed.

EP 0 325 262 A3



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number

EP 89 10 0913

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl. 4)
X	NATURE, vol. 331, no. 6151, 7th January 1988, pages 84-86, London, GB; A. TRAUNECKER et al.: "Soluble CD4 molecules neutralize human immunodeficiency virus type 1" * Whole article, especially figure 1 * ---	1-23	C 12 N 15/00 C 12 P 21/02 A 61 K 39/395 G 01 N 33/569// A 61 K 39/21
E	EP-A-0 314 317 (GENENTECH, INC.)(03-05-89) * Whole document * -----	1-23	
			TECHNICAL FIELDS SEARCHED (Int. Cl. 4)
			C 12 N
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 13-02-1990	Examiner CUPIDO M.
CATEGORY OF CITED DOCUMENTS X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons ----- & : member of the same patent family, corresponding document			